**Please note: this is the final draft of the accepted article:**

**Booth G, Williams D, Patel H, Gilbert AW. What is the content of virtually delivered pain management programmes for people with persistent musculoskeletal pain? A systematic review. In Press 2021; British Journal of Pain; Accepted 19th May 2021.**

**Please use the following link for the final, fully proofed and peer-reviewed journal article online:** [**https://journals.sagepub.com/loi/bjp**](https://journals.sagepub.com/loi/bjp)

# What is the content of virtually delivered pain management programmes for people with persistent musculoskeletal pain? A systematic review.

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# Abstract

**Introduction**

Virtual consultations (VC) have been embraced by healthcare organisations during the COVID-19 pandemic. VC allows continuation of patient care whilst adhering to government advised restrictions and social distancing measures. Multidisciplinary pain management programmes (PMP) are a core element of many pain services and utilising virtual methods to deliver PMPs has allowed them to continue to provide care. This systematic review aimed to explore the content of existing virtually delivered PMPs and discuss if and how these findings can be used to guide clinical delivery.

**Methods**

Eligible studies included adults (aged ≥18 years) with persistent musculoskeletal pain and any virtually delivered intervention that was described as a PMP or that had components of PMPs. Databases were searched from inception until July 2020. We performed a content analysis comparing existing interventions with established evidence-based clinical guidelines published by the British Pain Society (BPS). Intervention reporting quality was assessed using the Template for Intervention Description and Replication (TIDieR) checklist: an established checklist developed to improve the completeness of the reporting of interventions.

**Results**

Eight studies were included. One intervention included six of the seven components recommended by the BPS; none included all seven. “Skills training and activity management” was present in all eight interventions, “education” and “cognitive therapy methods” were present in six interventions, “graded activation” and “methods to enhance acceptance, mindfulness and psychological flexibility” were present in four interventions, “physical exercise” was present in two and “graded exposure” was present in one intervention. None of the studies described all 12 items of the TIDieR checklist adequately enough for replication.

**Conclusion**

Current virtual PMPs partially meet established clinical guidelines. Future virtual PMPs should be based on evidence-based clinical guidelines, and more research is needed to explore the effectiveness of virtually delivered PMPs and each recommended component.

**Keywords:**

COVID-19, chronic pain, persistent pain, pain management programme, rehabilitation, virtual consultations

# Introduction

The COVID-19 pandemic has led to significant disruption of services providing multidisciplinary pain management.1 The cessation of treatment for people with persistent pain will have consequences for service users and healthcare providers as the number of people requiring pain management services and the complexity of their problems will increase.2 Reduced access to assessment and treatment can lead to increased severity of symptoms2 which will add to the high prevalence of persistent pain across Europe.3 People with persistent pain are at more risk of being critically ill with COVID-19 and are more likely to be self-isolating in order to reduce their chances of being infected.4 Research has shown that people with persistent pain suffer greater adverse effects of ‘lockdown’.4 This may lead to self-perceived increases in pain, greater increases in anxiety and depression, increased loneliness and reduced physical activity.4 Consequences of reduced physical activity and greater stress increases the chance of physical deconditioning5 and risk of sarcopenia6, which is associated with a number of other chronic health conditions. Reduced access to multidisciplinary pain management services is likely to increase the burden on healthcare.6

Virtual consultations (VC) have been embraced by healthcare organisations in order to continue to provide patient care whilst complying with government advised social distancing measures.7 VC is one potential way to continue to provide pain services to patients during the pandemic2, 5 and has been demonstrated to be accurate, cost effective, safe8-12 and has high satisfaction among patients due to convenience, absence of travel time and reduced time off work.12.

Pain management programmes (PMP) consist of methods to promote behaviour change and enhance well-being.13 They aim to enable people with persistent pain to live as normal a life as possible by improving an individual’s ability to reduce and self-manage pain-associated disability and emotional distress.13 PMPs can be effective at reducing pain interference,14 pain intensity,15, 16 disability16 and distress.16 The British Pain Society (BPS) have published evidence-based clinical guidelines for pain management programmes in adults.13 There is currently no guidance for delivering PMPs virtually.

A recent rapid review17 presented a number of issues to consider when using VC to deliver group-based pain rehabilitation. These include issues with inclusion, access, participation, attrition, infrastructure, clinician training and confidence, and evaluation. Systematic reviews of the literature are important summaries of existing evidence to help inform changes to practice; to our knowledge, there are no systematic reviews discussing or evaluating virtually delivered PMPs. We therefore conducted a systematic review of the literature to identify studies reporting the use of VC in PMPs. This systematic review aims to explore the content of existing virtually delivered PMPs that include real-time delivery and discuss if and how these findings can be used to guide clinical delivery of virtual PMPs.

# Methods

This review has been reported in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement.18 The PRISMA Checklist can be seen in Supplementary Material 1. This review was registered with the International Prospective Register of Systematic Reviews (PROSPERO: CRD42020201593) on the 25th August 2020.

*2.1 Search strategy*

A list of search terms combining keywords and Medical Subject Headings (MeSH) terms was developed by the authors. Search terms related to chronic/persistent pain, pain management and telemedicine and were combined using Boolean operators. We searched four databases; CINAHL, MEDLINE, PsycINFO and AMED which were chosen due to their widespread coverage of clinical research topics. The search was completed by one author (AG) on 21st July 2020. We also searched the reference lists of similar systematic reviews for additional articles. See supplementary material 2 for the search strategy for each database.

*2.2 Study selection and eligibility criteria*

Search results were exported into Endnote19 and duplicates were removed. Remaining titles and abstracts were imported into Microsoft Excel for screening. Next, all titles and abstracts were screened against the eligibility criteria. One author (AG) screened 100% of the titles and abstracts, one author screened 50% (GB) and two authors screened 25% each (DW and HP) to ensure all titles and abstracts were independently screened by two authors. To ensure consistency, the eligibility criteria were discussed between all authors prior to starting the title and abstract screening to resolve any uncertainties. Furthermore, all authors referred to the eligibility criteria throughout and further uncertainties that arose were discussed with all authors at the nearest opportunity. Of the remaining articles, two authors (GB and AG) reviewed full-texts and agreed on the included articles. Any ambiguities in the screening process were resolved by discussion with a third author (DW).

Studies of interest were those that consisted of adults (aged ≥18 years) with persistent musculoskeletal (MSK) pain; defined as pain lasting ≥3 months,20 that is experienced in bones, joints, muscles, tendons and other related soft tissues.21 There were no restrictions to the specific conditions or origins of persistent MSK pain. Studies were included if they contained participants with persistent MSK pain, even if participants with other origins of pain were also included. We included any study with an intervention that was described as a PMP, a pain rehabilitation programme, or that had components that were consistent with PMP clinical guidelines,13 that was delivered virtually/remotely in real-time (e.g. via telemedicine). This included multi-component interventions as long as part of the intervention met this criterion. There were no limitations to study design. Studies of interest were published in peer-reviewed journals and had to be in English language. Papers reporting interventions where none of the components were aimed at active self-management of pain (such as prescription of medications/advice giving on medications) were excluded.

*2.3 Data extraction*

Three authors (GB, HP and DW) independently extracted the data using a data extraction table that was created based on the data sought in this study. One author (AG) reviewed all the data extraction for accuracy. Extracted data included study design, number of participants, participant characteristics (age, sex, ethnicity, conditions/pathologies), interventions characteristics (treatment approach, specific contents, providers, dose, method of delivery/technology used) and adverse events.

*2.4 Quality assessment (TIDieR)*

We completed a Template for Intervention Description and Replication (TIDieR) checklist22 for each study in order to report on the quality of the reporting of interventions. One author (GB) completed TIDieR checklists for all studies and a second author (AG) reviewed these for accuracy.

*2.5 Data synthesis*

We performed a content analysis of each intervention; comparing them with the “specific cognitive and behavioural methods” components of the established BPS PMP Guidelines for adults.13 These guidelines provide evidence-based recommendations of what PMPs delivered in healthcare settings should include. The guideline’s components include ‘graded activation’, ‘cognitive therapy methods’, ‘graded exposure’, ‘methods to enhance acceptance, mindfulness and psychological flexibility’, ‘skills training and activity management’, ‘physical exercise’, and ‘education’. Reference to the definitions used in the guidelines were referred to throughout this analysis to maximise accuracy.

# Results

*3.1 Study selection*

The search yielded a total of 1516 results, and 1155 unique articles were remaining after duplicates were removed. Of these, 1130 were excluded after title and abstract screening, leaving twenty-five articles for full text review. After reviewing the full texts, eight were included in the systematic review.23-30 The PRISMA flow diagram is demonstrated in Figure 1.

[INSERT FIGURE 1 AROUND HERE]

*3.2 Study and participant characteristics*

Of the eight included studies, one was a randomised controlled trial (RCT),24 one was a randomised trial (no control),23 two were non-randomised trials,25, 28 one was a longitudinal case-control study,26 One had a single-case experimental design,27 one was a retrospective cohort analysis29 and one was a descriptive study.30 Participant characteristics are demonstrated in Table 1.

*3.3 Intervention characteristics*

Table 1 presents intervention delivery methods, dose and schedule. All seven of the specific cognitive and behavioural methods of the BPS PMP guidelines13 were present across the studies. The highest number of guideline content in one study was six24 (table 2).

Table 1: Characteristics of included studies

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Study ref (First author, year) and design** | **No. of participants** | **Age: mean (SD) (years)** | **No. (%) Female** | **Conditions/ pathologies** | **Intervention details** | **Control/ comparator details** |
| Carmody, 2013  *Randomised trial* | Total:  101 | Total:  67.5 | Total:  3 (3%) | Not reported | Name:  Telephone-delivered cognitive behavioural therapy (T-CBT)  Delivery method:  Telephone  Individual  Dose:  12 sessions over 20 weeks  Schedule:  Weekly - first 8 sessions  Fortnightly – next 2 sessions  Monthly – final 2 sessions | Type:  Comparator  Description:  Telephone-delivered education (T-EDU) |
| Dear, 2015  *RCT (4-arm)* | Total:  490 | Total:  50 (13) | Total:  375 (80%) | Pain locations:  Head/facial  Neck  Shoulders  Upper limb  Lower back  Pelvic/sacrum  Lower limb | Name:  The Pain Course: internet delivered PMP based on CBT  Delivery method:  Online lessons  Lesson summaries  Homework tasks  Real-time contact: phone or email with a psychologist  No tailoring  Dose:  5 online lessons  5 lesson summaries  Real time:  *Regular contact* *= 10-15 mins weekly*  *Optional contact = 10-15 min sessions*  Schedule:  1 weeks between lessons 1 and 2, and 3 and 4  2 weeks between lessons 2 and 3, and 4 and 5 | Type:  Control  Description:  Waiting list |
| Gardner-Nix, 2012  *Non-randomised trial* | Total:  119 | Total:  52 (range 32 – 79) | Total:  90 (76%) | Back pain  Fibromyalgia  Facial pain  Arthritis  Neck pain  Shoulder pain  Lupus  Chest pain  Migraine  Multiple sclerosis  Neuropathy  Chrohn’s disease  Leg pain | Name:  Mindfulness based chronic pain management programme (MBCPM) – modified MBSR (stress reduction)  Delivery method:  Videoconferencing and face-to-face combination simultaneously  Class-sizes of 12-22 (combined on-site and off-site)  Dose:  12 sessions in 12 weeks  Schedule:  Weekly | Type:  Control  Description:  Waiting list |
| Helstrom, 2018  *Longitudinal case-control study* | Total:  160 | Total (all eligible participants):  76.7 (7.1) | Total (all eligible participants):  198 (79.2%) | High symptom chronic pain (pain interference 8+) | Name:  Pain Care Management (PCM)  Delivery method:  Telephone  Individual  Tailored  Dose:  20 min sessions  5 treatment sessions in 3 months  3 maintenance calls  Schedule:  Treatment: weeks 2, 4, 6, 9 and 12  Maintenance: weeks 16, 20 and 24 | Type:  Control  Description:  Symptom monitoring |
| Lafontaine, 2018  *Single-case experimental design* | Total:  2 | Total:  63 | Total:  2 (100%) | Fibromyalgia  Peripheral neuropathy | Name:  CBT for chronic pain  Delivery method:  Videoconference  Individual by a psychologist or doctoral student in psychology  Tailored  Dose:  10-13 1-hour sessions  Schedule:  Weekly | N/A |
| Mariano, 2019  *Non-randomised trial* | Total:  93 | Total:  57.1 | Total:  61 (65.6%) | Chronic low back pain (CLBP) | Name:  Group CBT teletherapy for chronic pain  Delivery method:  Online group CBT  Individual follow-up calls  8 per group  Not tailored  Dose:  8 sessions  Schedule:  Weekly | Type:  Control  Description:  In person group PMP |
| Mochari-Greenberger, 2017  *Retrospective cohort analysis* | Total:  170 | Total:  53.3 (8.9) | Total:  130 (76%) | MSK pain  Fibromyalgia  Nerve pain Headache | Name:  AbleTo - technology-enabled telebehavioural therapy program for chronic pain – CBT/ACT based  Delivery method:  Telephone or videoconference  Provider: behavioural health provider and behaviour coach  Individual  Tailored  Dose:  45 min sessions  15 sessions  8 weeks  Schedule:  Not reported | N/A |
| Palyo, 2012  *Descriptive study (qualitative)* | Not reported | Not reported | Not reported | Veterans with chronic pain | Name:  Cognitive-behavioural and physical therapy group intervention  Delivery method:  Videoconferencing  Group-based  Providers: psychologist and physical therapist  Dose:  2-hour sessions  10 weeks  Schedule:  Not reported | N/A |

*3.3.1 Graded activation guided by participant goals*

Graded activation guided by participant goals was present in four interventions.24, 26, 29, 30 Goal setting was used in all four of these studies and coincided with physical activity24, 26, 30 and self-care.29 “Relapse prevention”; a strategy aimed at facilitating maintenance of change, was present in two of these studies.24, 30

*3.3.2 Cognitive therapy methods*

Cognitive therapy methods were present in six studies.23, 24, 27-30 They were often described as teaching ways of managing or challenging negative thoughts.23, 24, 28, 29 Others simply stated they were using ‘cognitive restructuring’ methods27, 30without further elaboration of the detail of this.

*3.3.3 Graded exposure*

Graded exposure was described in one intervention.24 It was described as consisting of an explanation on the issues of fear and avoidance of physical activities, and instructions on gradually and safely increasing physical activities.

*3.3.4 Methods to enhance acceptance, mindfulness and psychological flexibility*

Four studies described methods of enhancing acceptance, mindfulness and psychological flexibility.24, 25, 28, 29 Two of these studies stated using mindfulness techniques,28, 29 one described monitoring negative thoughts,24 and one described exploring behaviours and responses to disability.25 There were no further explanations of these techniques.

*3.3.5 Skills training and activity management*

All eight studies included elements of skills training and activity management. Five studies included teaching of relaxation techniques24, 26-28, 30 to alter the adverse effects of thoughts and feelings. Another study stated teaching methods of managing negative emotions, without including specific details.23 Six studies taught goal setting.24, 26-30 Five studies taught activity management or pacing techniques24, 26-28, 30 to balance time spent on meaningful activity or rest. Four interventions included skills for communication and social interaction,23, 27, 28, 30 with two studies specifically stating teaching assertiveness skills.23, 30 Three studies were taught sleep management skills.25, 27, 28 One study was introduced to self-monitoring techniques.26 In addition to the two mentioned above, two other studies reported teaching “relapse prevention” skills.27, 28 These studies did not elaborate further on specifics of this skill.

*3.3.6 Physical exercise*

Two studies included physical exercise components.25, 30 One of these included stretching, strengthening and aerobic exercise30 and the other included training in exercise.25

*3.3.7 Education*

Six studies included education.24-28, 30 Three of these studies included pain education.24, 26, 27 The other studies included education on nutrition,25 weight management28 and posture.30

Table 2: Studies and items of the British Pain Society PMP Guidelines.

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| BPS PMP Content guidelines | Graded activation | Cognitive therapy methods | Graded exposure | Methods to enhance acceptance, mindfulness and psychological flexibility | Skills training and activity management (e.g. relaxation, pacing, sleep) | Physical exercise | Education (e.g. pain mechanisms/ physiology, pros and cons of medication) |
| Study (First author, year) |
| Carmody,  2013 |  | ✓ |  |  | ✓ |  |  |
| Dear,  2015 | ✓ | ✓ | ✓ | ✓ | ✓ |  | ✓ |
| Gardner-Nix, 2014 |  |  |  | ✓ | ✓ | ✓ | ✓ |
| Helstrom,  2017 | ✓ |  |  |  | ✓ |  | ✓ |
| Lafontaine,  2018 |  | ✓ |  |  | ✓ |  | ✓ |
| Mariano,  2019 |  | ✓ |  | ✓ | ✓ |  | ✓ |
| Mochari-Greenberger, 2017 | ✓ | ✓ |  | ✓ | ✓ |  |  |
| Palyo,  2012 | ✓ | ✓ |  |  | ✓ | ✓ | ✓ |
| ✓ = guideline item present in intervention. BPS = British Pain Society; PMP = pain management programme | | | | | | | |

*3.4 Control and comparator group characteristics*

Three studies had control groups; two of these studies24, 25 had waitlist controls and the other study compared online group CBT with in-person group CBT as the control.28 One study monitored symptoms and side effects of control participants.26 Another study had telephone delivered pain education as their comparator to telephone delivered CBT.23 Three studies did not have control or comparator groups due to their study design.27, 29, 30

*3.5 Intervention reporting quality (TIDieR)*

TIDieR checklist items one (brief name), two (rationale, theory or goal), three (materials), four (procedures), six (modes of delivery) and seven (locations) were adequately described in all eight intervention descriptions. Two intervention descriptions did not include sufficient information on who provided the intervention (item five).23, 28 One study did not adequately describe the dose or intensity (item eight)23 that the intervention was delivered at. Three studies did not provide information on whether the intervention was designed to be tailored or not, and under what circumstances (item nine).27, 28, 30 Seven studies did not mention details of any modifications (item 10) that may have happened during the study.23-26, 28-30 Five studies did not report any plans to measure adherence or fidelity (item 11)24-26, 28, 30 and five did not report any actual details of adherence or fidelity (item 12).25, 26, 28-30

One study adequately described seven of the TIDieR items,28 one study adequately described eight of the items,30 three studies adequately described nine items,23, 25, 26 two adequately described ten items24, 29 and one adequately described eleven items.27 No studies described all 12 items adequately enough for replication.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Study (First author, year) | Carmody, 2013 | Dear, 2015 | Gardner-Nix, 2014 | Helstrom, 2017 | Lafontaine, 2018 | Mariano, 2019 | Mochari-Greenberger, 2017 | Palyo, 2012 |
| TIDieR Checklist Item |
| 1. Brief name | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Why (rationale, theory, goal) | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Materials used in intervention (physical or informational) | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Procedures (activities, processes) | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Intervention providers | X | ✓ | ✓ | ✓ | ✓ | X | ✓ | ✓ |
| 1. Modes of delivery (e.g. face-to-face, virtual, individual or group) | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Location of intervention | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. When and how much (number of sessions, time period, schedule, dose) | X | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ | ✓ |
| 1. Details of tailoring | ✓ | ✓ | ✓ | ✓ | X | X | ✓ | X |
| 1. Modifications to intervention | X | X | X | X | ✓ | X | X | X |
| 1. Planned adherence and fidelity assessment | ✓ | X | X | X | ✓ | X |  | X |
| 1. Actual adherence and fidelity | ✓ | ✓ | X | X | ✓ | X | X | X |
| TIDieR = Template for Intervention Description and Replication checklist. ✓ = item adequately reported, X = item inadequately or not reported. | | | | | | | | |

Table 3: Studies and items of the TIDieR checklist

*3.6 Adverse events*

No studies formally reported any adverse events.

# Discussion

To our knowledge, this is the first systematic review to assess the content of virtually-delivered PMPs, or interventions including components of PMPs, in accordance with evidence-based clinical guidelines.13 We found that all eight of the included interventions had elements of ‘skills training and activity management’. ‘Cognitive therapy methods’ and ‘education’ were each present in six studies. ‘Graded activation guided by participant goals’ and ‘methods to enhance acceptance, mindfulness and psychological flexibility’ were each present in four studies. ‘Physical exercise’ was present in two studies and ‘graded exposure’ was only present in one study. None of the included studies included all the components recommended by the BPS guidelines.13 None of the studies adequately described all 12 TIDieR items: with details of modifications, fidelity and adherence being the items most commonly lacking. Name, rationale, theory or goal, materials, procedures, modes of delivery and location were adequately reported in all eight studies. No studies formally reported any adverse events.

None of the studies reported using clinical guidelines in their intervention design. ‘Cognitive therapy methods’ was one of the most common guideline items present across the interventions. Cognitive behavioural therapy (CBT) is effective (albeit effects are small or very small) at reducing pain and disability,16 and at improving multiple psychosocial issues associated with persistent pain.16, 31, 32 The effectiveness of CBT delivered by videoconference for posttraumatic stress disorder is comparable with face-to-face delivery.33 Psychotherapy methods delivered by videoconference, including CBT, can be effective for depressive symptoms34 and may be for anxiety;35 highly prevalent problems for people with persistent MSK pain conditions.36, 37 This suggests that CBT methods delivered by videoconference may be effective for people with persistent MSK pain and should be included in virtually delivered PMPs.

Patients with persistent non-cancer pain expect explanations or improved understanding of the their pain problem.38 Pain neuroscience education can facilitate a patient’s ability to cope with their pain, reduce kinesiophobia and decrease catastrophising.39 The evidence for patient education delivered via real-time virtual methods is lacking but may be feasible40 and should be considered for inclusion in virtual PMPs.

Half of the interventions included ‘methods to enhance acceptance, mindfulness and psychological flexibility’. Acceptance and mindfulness-based interventions can be effective at reducing pain and psychological problems and can improve physical and mental health-related quality of life in people with persistent pain.41, 42 Studies of stand-alone virtually delivered acceptance and/or mindfulness interventions are lacking in healthcare research; this lack of supporting evidence may be a reason these treatments were not utilised across all studies.

Physical exercise was reported in two studies. Exercise alone delivered via telemedicine has been shown to be better than no intervention.43 A combination of exercise and pain coping skills training delivered by videoconference is effective for pain and function long-term in people with chronic knee pain.44 Exercise should be included in virtual PMP's. All eight studies included ‘skills training’ which could be incorporated with exercise.

‘Graded exposure’ was present in only one study; this was taught during one online lesson and not practiced with the participants.24 Reasons for limited inclusion in interventions are unknown and, to our knowledge, there are no studies investigating graded exposure performed via virtual methods for any condition. Considering greater degrees of kinesiophobia are associated with higher pain levels and disability, and lower quality of life45 and the fact that graded exposure can reduce pain-related kinesiophobia,46 studies of virtually-delivered graded exposure are required to be able to evaluate their effectiveness when working remotely with patients.

*4.1 Strengths and limitations*

These results must be considered in light of the study limitations. There were multiple study designs included and only two of these were randomised trials. We were therefore unable to perform formal quality assessments of the included studies due to variation in the study types as we would be unable make useful comparisons.

This systematic review was prospectively registered on the PROSPERO database, it has been reported in line with PRISMA guidelines18 and multiple authors were involved at each stage of the review process. We compared existing interventions with established evidence-based clinical guidelines13 and we assessed intervention reporting quality with well-known guidance;22 strengthening our analysis. The included studies represented a broad range of chronic pain conditions, with a large number of these MSK in origin and location, although only one study solely focused on MSK pain.28

*4.2 Implications for research and clinical practice*

There is a current lack of high-quality research into real-time virtually delivered PMPs. There is also limited research into real-time virtual delivery of any of the recommended components of these programmes. Future research should address these gaps in order to improve confidence in either programme or individual component effectiveness, which can subsequently be used to guide evidence-based clinical practice. Future research should focus on delivering PMPs that are in-line with established evidence-based clinical guidelines such as the BPS PMP guidelines13. Interventions need to be well reported to enable replication in clinical practice; the use of checklists and tools such as the TIDieR checklist13 can aid this. Studies should also report any adverse effects of virtual PMPs as these are currently unknown, and patient safety remains of high importance. No specific guidance on clinical delivery can be made from this study. Despite the limited evidence for real-time virtual delivery of PMPs, clinical guidelines such as the BPS PMP guidelines13 should still be used as a framework when designing virtually delivered programmes, with requirements and resources being tailored to individual context.

# Conclusion

The COVID-19 pandemic has interrupted the care of people requiring multidisciplinary pain management for musculoskeletal pain. Virtually delivered PMPs have been cited as a means to continue the delivery of care for people with persistent pain. We have successfully conducted a systematic literature review of studies reporting virtually delivered PMPs. We found that few studies considered best practice guidelines, such as those offered by the British Pain Society during the design of their programmes. Clinical interventions should be based on evidence-based clinical guidelines and future research should explore the effectiveness of each component when delivered remotely.

**Authors Contributions**

GB and AG conceptualised and designed the study. GB and AG developed the search strategy. AG completed the search. GB, DW, HP and AG contributed to the screening process, data extraction. GB conducted the content analysis. All authors contributed to interpretation of results. GB drafted the manuscript. DW, HP and AG critically revised the manuscript. All authors approved the final version for publication.

**Funding**

This study was unfunded. Gregory Booth completed this work during an ICA Pre-doctoral Clinical Academic Fellowship supported by Health Education England and the National Institute for Health Research (Grant number: NIHR300342). Anthony Gilbert, Clinical Doctoral Research Fellow (ICA-CDRF-2017-03-025) is funded by Health Education England and the National Institute for Health Research (NIHR). The views expressed in this publication are those of the author(s) and not necessarily those of the NHS, the National Institute for Health Research or the Department of Health and Social Care.

**Conflict of interest**

The authors declare no conflict of interest.

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