Physical and functional outcomes following multidisciplinary residential rehabilitation for prearthritic hip pain among young active UK military personnel

Russell J Coppack,1,2 James L Bilzon,2 Andrew K Wills,3 Ian M McCurdie,4 Laura K Partridge,4 Alastair M Nicol,4 Alexander N Bennett1,5

ABSTRACT

Background: There are no studies describing the clinical outcomes of a residential, multidisciplinary team (MDT) rehabilitation intervention for patients with prearthritic hip pain. The aim of this cohort study was to describe the functional and physical outcomes of multidisciplinary residential rehabilitation for UK military personnel with prearthritic hip pain.

Methods: Participants (N=40) with a mean age of 33 years referred to a specialist residential rehabilitation centre completed a comprehensive multidisciplinary residential intervention. The main outcome measures were mean pain, physical function (modified shuttle test (MST) and Y-balance test), hip range of motion (HROM) and a patient-reported outcome measure (The Copenhagen Hip and Groin Outcome Score, HAGOS). All scores for symptomatic hips were taken at baseline and post-treatment.

Results: There were improvements in the Y-balance test and HROM following rehabilitation. There were significant improvements in mean difference (T1-to-T2) for Y-balance scores (15.8 cm, 95% CI 10.7 to 20.9, p<0.001), HROM (6.5° increase in hip flexion, 95% CI 4.6 to 9.4, p<0.001) and hip internal rotation (4.6°, 95% CI 2.7 to 6.6, p<0.001). Scores for HAGOS, pain, MST and functional activity assessment showed no improvement.

Conclusions: Among UK military personnel with prearthritic hip pain, MDT residential rehabilitation resulted in improvements in a functional Y-balance test, hip flexion and internal rotation. The study suggests short-term benefits across some outcomes for the current UK military approach to MDT residential rehabilitation.

INTRODUCTION

Prearthritic hip disorders are abnormalities of the articulating surfaces of the acetabulum and femoral head before the onset of osteoarthritis (OA), including intra-articular structures such as the acetabular labrum and chondral surfaces.1 Abnormalities of these structures can lead to biomechanical changes associated with significant hip pain and dysfunction in young adults.2,3 Femoroacetabular impingement (FAI) is recognised as a major cause of prearthritic hip pain in young active adults.4,5 FAI is characterised by impingement between the proximal end of the femur and acetabular rim caused by either an abnormally shaped femoral head (cam impingement) and/or an abnormally shaped acetabulum (pincer impingement).6,7 Abnormal femoroacetabular morphology is commonly seen in young men who make up much of the UK military population.8 Furthermore, hip pain with mechanical symptoms, particularly acetabular labral tears, are common symptoms presenting to the military orthopaedic surgeon.9

Hip arthroscopy and non-surgical interventions such as the treatments offered by physiotherapists, are recommended in the management of prearthritic FAI.1,10,11 However, while single (unidisciplinary) treatments have been the subject of research,2,10,11 there are no studies evaluating multidisciplinary team (MDT) programmes for the conservative treatment or postsurgical rehabilitation of prearthritic intra-articular...
hip disorders in young adults. Residential (inpatient) MDT rehabilitation for patients with musculoskeletal injuries has a long tradition in the UK military. Rehabilitation most often takes place in two clinical settings: outpatient primary care rehabilitation facilities (PCRFs), and inpatient residential centres including regional rehabilitation units (RRUs) and the UK Defence Medical Rehabilitation Centre (DMRC), Headley Court. Rehabilitation at the PCRF is focused primarily on physical function led by a physiotherapist, whereas residential centres have access to larger consultant-led MDTs delivering a broader biopsychosocial approach to treatment.

Despite the popularity and acceptance of residential MDT rehabilitation in the UK military, evidence supporting its effectiveness is lacking, and the extent to which outcomes differ between residential and outpatient approaches remains unknown. Therefore, the objective of our study was to describe the structure and process of a residential MDT intervention for UK military personnel with prearthritic hip pain and describe their associated clinical outcomes.

METHODS
Setting
The study was conducted at a specialist UK military rehabilitation centre.

Study design
We performed a retrospective cohort study. A review of the computerised clinical records was performed on military personnel admitted to the UK DMRC hip pain treatment programme. The intention of the observational study design was to describe and record the clinical outcomes of residential, multidisciplinary hip rehabilitation. Ethical approval was granted by the University of Bath Research Ethics Approval Committee for Health (REACH reference EP 15/16 87). In accordance with Ministry of Defence (MOD) research ethics guidelines, this study was exempt from requiring formal MOD ethical approval as it was categorised as a service evaluation. Approval for anonymised data extraction was granted in accordance with the local Caldicott guardian policy.

Participants
All UK military patients with clinical indicators of prearthritic, intra-articular hip pain attending DMRC for their first admission between October 2014 and May 2015 were included in the study. Clinical diagnoses were confirmed on imaging with MRI±MR arthrogram prior to admission for rehabilitation (figure 1). X-ray was used to identify typical FAI features including cam and pincer-type abnormalities. Patients were also included if they were admitted for treatment following arthroscopy for the management of acetabular labral tears. Participants comprised a mix of non-surgical and postsurgical patients completing concurrent residential rehabilitation reflecting current UK military rehabilitation practice.

Intervention
All patients completed an intensive, goal-based multidisciplinary residential rehabilitation programme. Patients received treatment from Monday to Friday for 5 h/day over a 3-week period (mean (SD) duration 16.2±1.3 treatment days). The MDT consists of a consultant physician, physiotherapist, exercise rehabilitation instructor and occupational therapist. On admission, all patients underwent a comprehensive clinical assessment with each member of the MDT. The standardised treatment goals were to improve hip range of motion (HROM), improve ‘global’ hip strength, improve core and trunk muscle function, improve strength and neuromotor control of the deep hip stabilizer muscles, correct gait and balance deficits, control/reduce pain, improve function in daily living, and weight management. Active therapies included individually prescribed and group exercises, manual physiotherapy techniques, hydrotherapy/swimming, and patient education. Administration of non-steroidal anti-inflammatory drugs (NSAIDs) and analgesics were minimised where possible and controlled by the supervising lead consultant. The programme was individually tailored to each patient based on the findings of repeat assessments, weight-bearing status and response to treatment. All patients received a home-based exercise programme on discharge. Table 1 summarises the treatment intervention.

Outcome measures
Relevant descriptive, sociodemographic and disease-related information at the time of admission were recorded. The primary outcome was the change from admission (baseline) to discharge (3 weeks) in mean scores on a series of standardised, validated outcome measures assessing pain, physical function and occupational employability. Hip and groin disability was measured using the Copenhagen Hip and Groin Outcome Score (HAGOS). This self-report questionnaire is a quantitative measure of hip disability containing 37 questions covering six domains. Each question is answered on a Likert scale and each subscale is transformed and scored separately on a 100-point scale where 0=extreme problems and 100=no problems. The HAGOS has good-to-excellent test–retest reliability and responsiveness to change, and is specifically designed to assess hip disorders in young active adults similar to the UK military population. Hip pain was scored using a 100 mm horizontal visual analogue scale (VAS) where 0 represents no pain and 100 worst possible pain. A minimal clinically important difference is 18 mm. The VAS has shown good internal consistency and has been extensively evaluated in clinical trials.

Active HROM was measured for flexion and internal rotation using a 360° goniometer. This is a standard assessment widely used in the clinical setting with high

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test-retest reliability values reported in participants with hip OA.18 19

Physical performance was measured using the modified 20 m shuttle test (MST). The MST20 involves walking and running at gradually increasing speeds until the patient is unable to continue. Speed is controlled by a series of paced-auditory cues accompanied by recorded verbal instructions. The MST ranges from level 0 (0.66 m/s) to level 28 (5 m/s). Thirty shuttles at level 28 is the highest level attainable before the test is terminated. The MST is preferred by military rehabilitation practitioners to the 6 min walk test21 because the incremental increase in speed mimics a military marching task.22 Dynamic balance and postural control was measured using the modified star excursion balance test, known as the Y-balance test (figure 2).23 The Y-balance test was developed to incorporate those directional movements (anterior, posteromedial, posterolateral directions) with the greatest accuracy in identifying lower extremity dysfunction.24 The supervising therapist measures the maximum distance in centimetres the patient can reach with the free limb while balancing on the contralateral limb. Scores for all three movements were combined to provide a single composite measure of dynamic balance. The test is a highly reliable tool to measure dynamic balance,25 and is currently used by the UK military.
Occupational employability was assessed using the functional activity assessment (FAA). The FAA is a single-item, five-point ordinal measure developed to assess occupational outcome in military rehabilitation (Table 2). The validity of the patient-reported FAA has been tested on our military patient population at DMRC and reported in the scientific literature. All outcome measures were recorded at baseline (pretreatment) and at 3 weeks (post-treatment) by a trained member of the MDT.

### Table 1 Multidisciplinary team residential hip rehabilitation programme—components of treatment

<table>
<thead>
<tr>
<th>Treatment modality</th>
<th>Treatment content</th>
<th>Treatment goals</th>
<th>Frequency per week (duration)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group exercise</td>
<td>Strengthening exercises, active range of motion exercises, functional balance drills, gait drills, progressive coordination drills, non-weight-bearing aerobic/ endurance exercise, minor team games</td>
<td>Restore strength of deep hip stabilisers, improve core strength, increase joint range of motion, improve balance and neuromotor control, improve muscle endurance, promote group cohesion and social support</td>
<td>12 (30–45 min)</td>
</tr>
<tr>
<td>Individual physiotherapy*</td>
<td>Manual therapy techniques, muscle activation and timing patterns, active and passive range of motion exercises, advice on home exercise, gait re-education training</td>
<td>Improve quality and timing of movement, improve muscle strength, reduce pain, increase joint range of motion, induce relaxation, promote normal walking gait</td>
<td>5 (30 min)</td>
</tr>
<tr>
<td>Hydrotherapy/swimming</td>
<td>Non-weight-bearing aerobic exercise, strengthening exercises, active range of motion exercises, self-paced recreational swimming, progressive/assisted weight-bearing exercise and activity</td>
<td>Improve muscle strength, improve aerobic capacity, increase joint range of motion, improve confidence in weight bearing, induce relaxation, promote enjoyment and variety of treatment</td>
<td>3 (60 min)</td>
</tr>
<tr>
<td>Individual occupational therapy†</td>
<td>Relaxation techniques, postural re-education, cognitive–behavioural therapy techniques, self-help coping strategies, pain management</td>
<td>Induce relaxation, promote behavioural change, control pain, correct/improve poor posture</td>
<td>3 (60 min)</td>
</tr>
<tr>
<td>Patient education</td>
<td>Coping with pain, benefits of exercise, joint protection, anatomy and pathology of hip pain, nutrition</td>
<td>Activity modification, reduction of pain, promote behavioural change, weight management, improve knowledge of treatment options, improve ability to relax, improve knowledge of self-help techniques</td>
<td>2 (60 min)</td>
</tr>
</tbody>
</table>

*Exercise dosage, progression and intensity were governed by the physiotherapist and tailored to the needs of each individual patient. †Occupational therapy referrals were individually prescribed to selected patients.

### Figure 2 Y-balance test. From a single-leg stance the participant reaches the freely moveable limb along a line in the anterior (A), posterolateral (B) and posteromedial (C) directions.
score/2) to provide an aggregate score for internal rotation, flexion and dynamic balance pre-to-post treatment. All continuous variables were assessed using the Kolmogorov-Smirnov test and an examination of histograms and their skewness and kurtosis. None were considered to deviate from normality. Descriptive statistics were calculated for the whole sample. Continuous variables are presented as means and SD, and categorical variables are presented by frequencies and percentages. Within group changes over time were analysed using a paired t test. Ordinal data for the FAA employability scale were assessed using the Wilcoxon signed rank test. Statistical calculations were performed using SPSS (V.21.0.0, SPSS Inc, Chicago, Illinois, USA).

RESULTS

Participant characteristics
The demographic and baseline patient characteristics are presented in table 3. Forty UK military patients (27 (67.5%) male) with a mean age of 32.8 years (SD 7.1), range 20–50 years, were included in the study. The distribution of patients by military branch was 26 (65%) Army, 10 (25%) Royal Air Force (RAF) and 4 (10%) Royal Marines (RM). The patient distribution by rank seniority was 23 (57.5%) junior ranks, 10 (25%) senior ranks and 7 (17.5%) officer rank. Nineteen (47.5%) patients had undergone arthroscopic repair for a labral tear, and 29 (72.5%) had completed a previous course of inpatient treatment at a RRU.

Main outcome measures
The main treatment effects (paired t test) for all outcome measures are presented in table 4.

Physical and functional outcome measures (Y-balance, HROM, MST)
The change in Y-balance test score over time was statistically significant. The mean (SD) pretreatment and post-treatment scores for Y-balance were 240.5 (26.9) cm and 256.3 (20.8) cm, respectively. The mean improvements of 15.8 cm showed a significant increase (p<0.001) in Y-balance score (95% CI 10.7 to 20.9). The HROM scores (flexion and internal rotation) also revealed significant improvements over the period of treatment. The mean (SD) scores for flexion improved by 6.5° (95% CI 4.6 to 9.4, p<0.001) from 110.2° (24.3) at baseline to 116.7° (23.3) at week 3. The mean (SD) scores for internal rotation improved by 4.6° (95% CI 2.7 to 6.6, p<0.001) from 25.2° (13.7) at baseline to 29.8° (12.4) at week 3. There were no pre-to-post treatment differences in the MST scores. The mean (SD) test scores achieved were level 10.3 (4.9) at baseline and level 10.5 (5.1) at week 3. The small mean difference of 0.2 levels was not statistically different from baseline (95% CI –1.2 to 0.7, p=0.59).

Patient-reported outcomes (pain, HAGOS)
The pre-to-post treatment scores for changes in pain were not significant. The baseline mean (SD) VAS was 36.3 (24.2) mm and post-treatment score was 34.8 (22.3) mm. The mean difference was a 1.5 mm reduction in pain after treatment (95% CI –0.49 to 8.0, p=0.63). There were no significant pre-to-post treatment improvements in scores on any of the HAGOS subscales. Table 4 and figure 3 shows that five out of six subscale domains deteriorated over the period of treatment based on the patients self-ratings.

Occupational employability (FAA)
The median score on the FAA occupational employability index did not change from pretreatment (Md=3.0) to post-treatment (Md=3.0), p=0.13. There were no between-group differences for any outcome measure in the surgical versus non-surgical, or male versus female subgroups.

DISCUSSION
In this first study of a short-term MDT residential rehabilitation intervention in military personnel with...
prearthritic hip pain, we report improved HROM and dynamic balance test scores. There was no evidence for a change in outcome scores for the patient-reported HAGOS subscales, pain, MST and FAA employability scale.

How do these results compare with other studies?

The results from our study are inconsistent with the findings of two recent publications reporting HROM, pain and functional outcomes following conservative treatment of prearthritic hip disorders. Both studies employed a within-group observational design similar to the present study, and a comparable population in terms of sample size, mean age, diagnostic inclusion criteria, and baseline pain-related and function-related impairment. Fundamental to all three programmes was an exercise-based regimen augmented by education, advice on activity modification and manual physiotherapy. Hunt et al. found significant improvements in pain (numeric pain scale), physical function (Western Ontario and McMaster Universities Arthritis (WOMAC) index; non-arthritic hip score (NAHS)) and quality of life (Short Form-12) in patients with prearthritic, intra-articular hip pain completing a directed course of conservative treatment. Emara et al. showed a stage-based physical therapy programme-improved pain symptoms and patient-reported function (Modified Harris Hip Score (MHHS)), but had no effect on HROM, in 33 patients with FAI up to 28 months after treatment. This contradicts our findings showing significant post-treatment improvement in patients HROM (minimal clinically important difference, 5°), with no change in pain or patient-reported function measured by the HAGOS.

Between-study differences in outcome measures (MHHS; NAHS; WOMAC; HAGOS), time to follow-up (3 weeks; 12 months; 28 months) and residential versus outpatient settings may explain the inconsistencies in results. The observational design of our study limits the conclusions that can be drawn concerning the causal effects of treatment on rehabilitation outcomes. Further high-quality randomised trials investigating longer term effects of rehabilitation with our military population are required. This is particularly important as there may be a delayed period of adaptation before reductions in pain

### Table 3 Baseline descriptive characteristics of study participants (N=40)

<table>
<thead>
<tr>
<th>Variable/characteristic</th>
<th>Male (n=27)</th>
<th>Female (n=13)</th>
<th>Total (n=40)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, year, mean (SD)</td>
<td>32.8 (7.1)</td>
<td>32.9 (3.8)</td>
<td>32.8 (7.1)</td>
</tr>
<tr>
<td>Height, cm, mean (SD)</td>
<td>178.4 (5.3)</td>
<td>169.3 (3.2)</td>
<td>175.4 (6.4)</td>
</tr>
<tr>
<td>Weight, kg, mean (SD)</td>
<td>81.8 (11.5)</td>
<td>75.9 (16.9)</td>
<td>79.7 (13.5)</td>
</tr>
<tr>
<td>Body mass index, kg/m², mean (SD)</td>
<td>25.5 (3.5)</td>
<td>26.5 (6.1)</td>
<td>25.8 (4.5)</td>
</tr>
<tr>
<td>Treatment duration, days, mean (SD)</td>
<td>16.2 (1.3)</td>
<td>16.1 (1.2)</td>
<td>16.2 (1.3)</td>
</tr>
<tr>
<td>Service branch, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Army</td>
<td>18 (66.7)</td>
<td>8 (61.5)</td>
<td>26 (65.0)</td>
</tr>
<tr>
<td>RAF</td>
<td>5 (18.5)</td>
<td>5 (38.5)</td>
<td>10 (25.0)</td>
</tr>
<tr>
<td>RM</td>
<td>4 (14.8)</td>
<td>0 (0.0)</td>
<td>4 (10.0)</td>
</tr>
<tr>
<td>Rank seniority, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Junior rank</td>
<td>15 (55.5)</td>
<td>8 (61.5)</td>
<td>23 (57.5)</td>
</tr>
<tr>
<td>Senior rank</td>
<td>8 (29.7)</td>
<td>2 (15.5)</td>
<td>10 (25.0)</td>
</tr>
<tr>
<td>Officer rank</td>
<td>4 (14.8)</td>
<td>3 (23.0)</td>
<td>7 (17.5)</td>
</tr>
<tr>
<td>Diagnosis, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FAI/acetabular labral tear (right)</td>
<td>7 (25.9)</td>
<td>6 (46.2)</td>
<td>13 (32.5)</td>
</tr>
<tr>
<td>FAI/acetabular labral tear (left)</td>
<td>7 (25.9)</td>
<td>3 (23.0)</td>
<td>10 (25.0)</td>
</tr>
<tr>
<td>FAI/acetabular labral tear (bilateral)</td>
<td>3 (11.1)</td>
<td>3 (23.0)</td>
<td>6 (15.0)</td>
</tr>
<tr>
<td>Other hip pain diagnosis</td>
<td>10 (37.0)</td>
<td>1 (7.8)</td>
<td>11 (27.5)</td>
</tr>
<tr>
<td>Previous surgery, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arthroscopy</td>
<td>7 (26.0)</td>
<td>12 (92.3)</td>
<td>19 (47.5)</td>
</tr>
<tr>
<td>‘Other’ surgery</td>
<td>7 (26.0)</td>
<td>0 (0.0)</td>
<td>7 (17.5)</td>
</tr>
<tr>
<td>No surgery</td>
<td>13 (48.1)</td>
<td>1 (7.7)</td>
<td>14 (35.0)</td>
</tr>
<tr>
<td>Previous treatment, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RRU</td>
<td>Yes</td>
<td>21 (77.7)</td>
<td>29 (72.5)</td>
</tr>
<tr>
<td>PCRF</td>
<td>No</td>
<td>6 (22.2)</td>
<td>11 (27.5)</td>
</tr>
<tr>
<td>Medication, N (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>25 (92.6)</td>
<td>11 (84.6)</td>
<td>36 (90.0)</td>
</tr>
<tr>
<td>No</td>
<td>2 (7.4)</td>
<td>2 (15.4)</td>
<td>4 (10.0)</td>
</tr>
<tr>
<td>Treatment duration, days, mean (SD)</td>
<td>16.2 (1.3)</td>
<td>16.1 (1.2)</td>
<td>16.2 (1.3)</td>
</tr>
</tbody>
</table>

FAI, femoroacetabular impingement; PCRF, primary care rehabilitation facility; RAF, Royal Air Force; RM, Royal Marines; RRU, regional rehabilitation unit.
are realised following 3 weeks of intensive rehabilitation, which appears to have stimulated improvements in dynamic balance and HROM.

When considering the structure and process of rehabilitation, comparisons between our study and other similar studies are extremely limited due to the lack of published research focusing on MDT residential treatment for prearthritic hip pain. In terms of structure, the most relevant findings are reported in patients with hip OA. Angst et al found a comprehensive 3-week residential rehabilitation intervention led to statistically and clinically important improvements in pain and function (WOMAC) for patients with comorbid hip pain. In an older study, Weigl et al reported improvements in pain and physical function at 2-year follow-up in 44 patients with hip OA completing a 3–4-week residential programme.

The duration and content of treatment in our study closely approximates those utilised by Angst and colleagues and Weigl and colleagues (eg, 3 weeks group exercise, individual physiotherapy, patient education, NSAIDs); however, we did not observe similar improvements in patients’ pain and function following residential treatment. The composition of the MDT, study population and treatment outcomes differed between studies, and a potential reason for the discrepancies between our findings and those of previous studies is the heterogeneity with respect to participant case mix, outcome measures, treatment setting (hospital vs specialist rehabilitation centre) and MDT size and composition. These methodological inconsistencies must be addressed to allow a better understanding of the benefits of MDT residential rehabilitation for prearthritic hip pain.

Patient-reported outcomes (PROs) are considered the gold standard when measuring the patient’s perspective of treatment efficacy. The HAGOS did not demonstrate any significant pre-to-post treatment improvements in our study. This PRO is used in the UK military hip rehabilitation pathway because it was developed for physically active young-to-middle-aged adults with hip and/or groin pain and is designed to assess treatment-induced changes from week-to-week. However, while the HAGOS subscales have shown good test-retest reliability and responsiveness in athletic populations, its performance in a military population is unknown, and it is possible this scoring system fails to address activities of most relevance to Armed Forces personnel. Studies reporting clinically meaningful changes in other PROs following hip rehabilitation used a minimum 6-week period between tests. The reference values of the HAGOS over longer testing periods with a military population need to be established.

The primary aim of UK military rehabilitation is to return personnel fit to undertake their technical trade and general duties. Therefore, outcome measures providing information on a patient’s military-specific occupational status are important. The FAA scores showed the same median rating of 3.0 (unfit for trade but fit for

### Table 4 Pre-to-post rehabilitation mean differences (paired t test) for all outcomes

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Pretreatment mean (SD)</th>
<th>Post-treatment mean (SD)</th>
<th>Difference, mean (95% CI)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y-balance*</td>
<td>240.5 (26.9)</td>
<td>256.3 (20.8)</td>
<td>15.8 (10.7 to 20.9)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HROM (flexion)*</td>
<td>110.2 (24.3)</td>
<td>116.7 (23.3)</td>
<td>6.5 (4.6 to 9.4)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HROM (internal rot)*</td>
<td>25.2 (13.7)</td>
<td>29.8 (12.4)</td>
<td>4.6 (2.7 to 6.6)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Modified shuttle test</td>
<td>10.3 (4.9)</td>
<td>10.5 (5.1)</td>
<td>0.2 (–1.2 to 0.7)</td>
<td>0.60</td>
</tr>
<tr>
<td>Pain (VAS 1–100 mm)</td>
<td>36.3 (24.2)</td>
<td>34.8 (22.8)</td>
<td>1.5 (–0.5 to 8.0)</td>
<td>0.63</td>
</tr>
</tbody>
</table>

†A normalised score (100 indicating no symptoms and 0 indicating extreme symptoms) is calculated for each subscale.

ADL, activities of daily living; HAGOS, Copenhagen Hip and Groin Outcome Score; HROM, hip range of motion; PA, participation in physical activity; QOL, quality of life; VAS, visual analogue scale.

**Figure 3** Participants (N=40) mean Copenhagen Hip and Groin Outcome Score (HAGOS) pre-to-post treatment. Total score for each subscale is summed and transformed such that 100=best score (no problems) and 0=worst score (extreme problems). Data are presented as mean (95% CI).
restricted general or military duties) before and after treatment. The FAA has shown adequate construct and concurrent validity as a surrogate measure of physical health; however, our results suggest its ability to track changes in employment status following a single residential admission period is unproven.

Study limitations

Despite the novelty of these findings, they must be considered in the light of limitations of the study design. It is a retrospective cohort study with no control group. This limits the conclusions that can be drawn on the effectiveness of our rehabilitation programme. Although improvement in symptoms and function could be a result of treatment, a control group is needed to gain a causal estimate of the effect of rehabilitation. Prospective studies employing a randomised controlled design should be undertaken to compare available treatment options. The lack of follow-up beyond 3 weeks would not capture any longer term benefits of rehabilitation which may explain the non-significant findings for some outcomes in our study. Therefore, we have restricted our comments to the 3-week period of rehabilitation and do not speculate on the long-term benefits of MDT residential rehabilitation. Future studies should address longer term compliance to and effectiveness of conservative treatment. While the use of a well-defined military population may limit the generalisability of our results to other populations and settings, we believe the findings may also be relevant to young active sportsmen and women. Our patients underwent rehabilitation at varying stages of their recovery, and while this approach mimics our clinical practice, some heterogeneity in clinical severity of the sample may have attenuated the treatment effect. Finally, we did not record psychological variables (e.g., anxiety, irritability, depression) that may be related to pain. Notwithstanding these limitations, the observational design did provide the opportunity to obtain data with a young active cohort, and document the acute responses to treatment reflecting the clinical reality of a military rehabilitation setting.

CONCLUSIONS

With increased pressure to ensure military personnel remain fit for operations against the backdrop of reductions in manpower, it is expected that intensive residential rehabilitation will become an increasingly important component of future healthcare provision in the UK Armed Forces. This is particularly important given recent increases in operational deployments and the commensurate increase in occupational exposure associated with loaded marching. This study is the first to report outcomes of MDT residential rehabilitation in a military cohort with prearthritic hip pain.

Comprehensive residential treatment showed evidence of improvements in a functional capacity (Y-balance) test and HROM (flexion, internal rotation). The intervention did not confer any benefits on pain, the patient-reported HAGOS, or occupational employability (FAA). The study is strongly suggestive of some short-term benefits for the current UK military approach to MDT residential rehabilitation in the management of prearthritic hip pain. Further studies should adopt longer follow-up observations, evaluating longer term adaptations and assessing compliance to post-residential treatment plans. There is a clear need for further research using randomised designs examining MDT residential rehabilitation against other treatment options to ensure patients receive treatment in the optimal clinical setting. Future studies should also focus on the measurement properties of hip PROs with a military population.

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Contributors RJC designed the study, conducted the initial analysis, drafted the initial manuscript and approved the final manuscript as submitted. JLB, AKW and ANB supervised the conduct of the study, assisted with data analysis, revised and approved the manuscript, and approved the final manuscript as submitted. IMM, AMN and LKP collected the data, reviewed and revised the manuscript, and approved the final manuscript as submitted.

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