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Research Article

The Effectiveness of a 12-week Exercise Programme in Hip OsteoArthritis: A Randomised Controlled Trial

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Abstract

Purpose: Osteoarthritis (OA) is a common disease caused by the destruction of the joint cartilage. Pain, stiffness and dysfunction during activities of daily living are common reasons leading to a hip arthroplasty. Programmes based on quadriceps strengthening are now evidence-based. Therefore, this study aims to explore and investigate the effects of a 12-week hip exercise programme on pain and functional levels.

Methods: This study reports outcomes of a randomised controlled trial. 40 participants were randomly allocated into 2 groups: control group and an intervention group. All participants were asked general demographic data at baseline, followed by assessment using the NPRS, WOMAC Scale, 6 MWT and muscle testing. Participants in the control group received no intervention, whereas the others performed hip exercises. A 12-week exercise programme resulted in highly statistically significant improvements for participants in the intervention group and minimal changes for the control group.

Results: Data is presented for both groups respectively as follows: 6MWD P<0.001, P=0.928; NPRS P=0.035, P=0.325; WOMAC P<0.001, P=0.032.

Conclusion: A 12-week exercise programme resulted in clinical and statistically significant improvements for the intervention group.

Key Words: Arthritis; Management; Physiotherapy; Rehabilitation

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Introduction

Osteoarthritis (OA) is one of the most common joint disorders worldwide [1,2], a condition which is responsible for pain, stiffness, and dysfunction during activities of daily living. One of the functional capacities affected by OA is deterioration in muscle strength mainly of the quadriceps, hamstrings, and other musculature around the hip [3-5].

An overwhelming amount of literature is available supporting the use of exercise to help manage or decrease joint pain as a result of knee OA, programmes which including muscle strengthening, aerobic/cardiovascular exercise and more [6-12]. Evidence based research about management through exercise of Hip OA is though limited. So as to counteract the limited information, guidance through recommendations for the non-pharmacological management of Hip OA were issued by the European League Against Rheumatism (EU-LAR) Osteoarthritis Task Force stating these patients should receive tailored programmes which include both long and short-term goals or action plans, regular evaluation by healthcare professionals and follow-up with the possibility of adjusting the programme accordingly [13]. These recommendation have been further supported by more recent guidelines issued by the American Physical Therapy Association 2017 [14]. the Ottawa Panel Clinical Practice Guidelines [15-17]. which although specific to knee OA, make reference to the hip, and the Osteoarthritis Research Society International (OARSI) guidelines, 2019 [18]. The latter set recommendations for flexibility, strengthening and endurance exercises aimed at addressing impairments in hip ROM, muscle weaknesses and reduced flexibility. These guidelines also recommend that for the dosage and duration to be effective, the exercises should be carried out 1-5 times per week over a 6-12 week period but this is still inconclusive and the duration still varies from as little as 6 weeks to up to 4months.

Results from studies offer varied outcomes. In a randomised trial by Bieler et al [10], 4 months of either physiotherapist-supervised, moderate, progressive, strength training (n = 50), physiotherapistsupervised Nordic Walking (n = 50), or unsupervised Home-Based Exercises (n = 52) showed no differences in muscle strength and active hip ROM with the authors suggesting that the patients recruited might not have been able to achieve the target load prescribed and that muscle strength and functional performance might not be correlated.

Svege et al [18] evaluated the effects of exercise therapy and patient education in a 12-week exercise programme, one which resulted in no beneficial long-term effects except for the significantly reduced pain levels reported during the 6-minute walk test at the 10th and 29th month. Svege et at [18] claimed that their participants had mild functional limitations at baseline, adherence to the programme was reported in 53% of the participants only and that the exercise programme may have been ineffective in dosage progression and execution of the exercises.

In contrast to the above, a 10-week programme of progressive explosive-type resistance training resulted in significant improvements in self-reported outcomes and increased lower limb muscle power [19]. This study reported significant improvements in self-reported ADL function, as well as leg muscle power.

Despite the vast amount of literature available on exercise in the management of most knee OA, research on the management of Hip OA is limited and no consensus regarding the optimal time-frame and dosage has yet been established. Therefore the authors aimed at investigating outcomes of a 12-week exercise programme in patients with Hip OA.

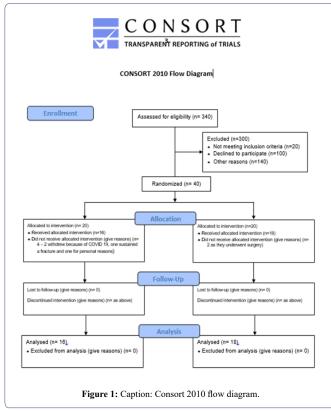
Methodology

Experimental approach

This paper reports a randomised controlled trial. Data obtained from both groups were recorded at the start of week 0 and the end of week 12. By adopting this design, a pre and post test method of data collection was implemented with all participants undergoing a series of outcome measures at the start of the study as described below and retested on all the participants upon completion of a 12-week intervention.

Subjects

All 340 participants who were waiting for a total hip replacement at the local hospital were eligible to participate. A total of 40 patients accepted to participate. In order to be recruited, participants had to be aged 40 years or over. Participants whom on assessments were found to have unstable cardiovascular or respiratory disorders, had recent lower limb surgery, had a condition affecting lower limb function, were unable to mobilise or were not able to commit to the exercise programme were excluded from the study. Procedures of recruited are outlines in Figure 1.



Procedures

At baseline, all participants were asked general demographic data including age, weight, height, body mass index (BMI), gender and drug history. The following outcome measures were then carried out on every subject at baseline (Week 0) and repeated at week 12.

Numerical pain rating scale (NPRS)

The NPRS is a widely used in the clinical setting as in research [20]. It is a simple and valid tool to use with acceptable construct validity for this tool (P < 0.001).

Western Ontario and McMaster universities scale and the 6 minute walking test

Functional Levels were assessed using the Western Ontario and McMaster Universities (WOMAC) Scale and a 6 MWT. The WOM-AC is a widely used instrument which is specific to assessing patients with hip and knee OA [21]. This assessment tool was self-administered. It consists of 33 items which evaluate the health and function of the patient from various aspects including clinical symptoms (5 questions), severity of joint stiffness (2 questions), degree of pain (9 questions), and activity of daily living (17 questions).

The six-minute walk test (6MWT) measures the distance an individual is able to walk over a total of 6minutes on a hard, flat surface. The individual was allowed to self-pace and rest as needed as they traverse back and forth along a marked walkway. Such a test has been reported to be a valid and reliable indicator of aerobic fitness [22]. The subjects were instructed that they could stop and rest during the test, but the time kept running. At the end of the test, the distance covered was recorded.

Muscle torque testing

All subjects also underwent peak isometric lower limb muscle torque testing for the following muscle groups: Quadriceps, Hamstrings, Adductors and Abductors using the Lafayette Hand held dynamometer. A maximum of three attempts were performed with the average result taken. Participants were instructed to exert maximal effort and were verbally coached to push as hard as they could for each recorded trial.

All patients were randomly assigned using a random computer generator to one of 2 groups by the intermediary: the control group who were awaiting surgery and were not receiving any regular physiotherapy and the Intervention group, who were subject to a set of hip exercises based on a protocol described by Deyle et al [23]. These exercises were carried out 3 times weekly for a period of 3 months. During the first 4weeks, all sessions were supervised by a physiotherapist, independent to this study who monitored and increased the duration or difficulty of exercises according to patients' progression. From the 5th to the 8th week, patients were requested to attend 2 supervised sessions and complete the third one at home. Between the 9th and the 12th week, patients attended one supervised intervention and completed 2 exercise sessions at home. All patients were given a copy of the exercise sheet with instructions for use at home.

Statistical analysis

Data was analysed using the statistical package for social science (SPSS) software version 27 to determine any significant changes over time between both groups using the ANOVA test. Comparision of outcomes at the 12th week between both groups was carried out using the Independent Sample's T-test. Effect sizes were calculated to identify the extent of the experimental effect using an online calculator:

https://www.socscistatistics.com/effectsize/default3.aspx.

Ethical approval was also obtained from the University Faculty Research Ethics Committee and registered with clinicaltrials.gov, having identifier number NCT04771936.

Results

The intervention lasted 12 weeks, with 34 patients completing the study. In the control group 2 participants were excluded from the study as they underwent a hip arthroplasty. In the intervention group, 4 participants withdrew for the following reasons: one participant sustained a fracture of her ribs, two participants quit attending because of an increase in Covid-19 cases and one participant dropped for personal reasons. Table 1 outlines the baseline characteristics of all the participants of both groups.

Baseline characteristics n= 51	Mean Value (SD)
Gender	Males - 28 Females - 23
Age (years)	60 years or less – 16 61-69 years – 21 70 years or more - 14
Affected hip	Right hip – 25 Left hip -21 Bilateral hips affected - 5
Weight (kg)	83.65 (+16.55)
Height (cm)	161.79 (+ 9.92)
BMI (kg/cm2)	31.96 (+5.89)
6MWT (m)	267.82 (+ 108.34)
NPRS	5.31 (+ 2.59)
WOMAC	49.63 (+ 19.69)
Core Stability	Good - 18 Poor - 33

N= number of participants; SD= standard Deviation; kg= kilograms; cm= centimetres; BMI= Body Mass Index; 6MWT= 6 Minute Walk Test; NPRS= Numerical Pain Rating Scale; WOMAC = Western Ontario and McMaster Universities Osteoarthritis Index

Outcome measures

Six Minute walk distance (6MWD)

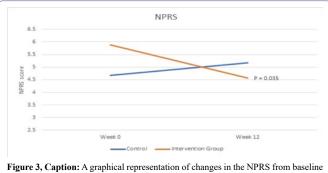
As can be noted in Figure 2, significant improvements in the 6 minute walking distance were recorded for participants in the intervention group. After 12 weeks of rehabilitation, a statistically significant increase in walking distance of 43m from a mean baseline value of 262.63m (SD 118.27) to 305.69m (SD 131.42), P< 0.001 (ES d=0.34) was obtained. Such improvements were not noted in participants of the control group. These participants registered no difference in walking distance from a mean baseline value of 257.94m (SD 127.80) at week 0 to a mean walking distance of 256.94m (SD 137.81) at week 12, p=0.928 (ES d=0.01).

Numerical pain rating scale (NPRS)

As is demonstrated in Figure 3, participants of the intervention group registered a significant decrease in pain levels of P=0.035, ES d=0.49 (mean baseline NPRS score of 5.88 [SD 2.68] at week 0 to 4.56 [SD 2.71] at week 12). Once again, such positive changes were not observed in participants of the control group, who registered an increase in pain values of 11% from their baseline (NPRS at week 0 4.67 [SD 2.87] to 5.17 [SD 2.92] at week 12), P=0.325, ES d= 0.17.



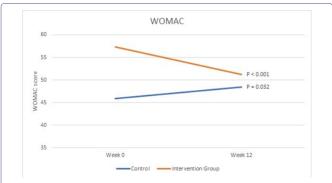
Figure 2. Caption: A graphical representation of changes in the 6MWD from baseline to week 12. Statistically significant changes are noted in the graph, through the representation of the p-value. N.B. 6MWD (m)= Six-minute walk distance measured in metres.

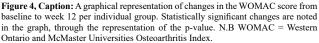


to week 12. Statistically significant changes are noted in the graph, through the representation of the p-value. N.B. NPRS= Numerical Pain Rating Scale.

Western ontario McMaster universities osteoarthritis index (WOMAC)

Following a 12-week exercise intervention, a significant improvement in the impact of OA in participants forming part of the intervention group of 11%, P<0.001, ES d=0.32 resulted. This group scored a value of 57.31 (SD 18.39) at week 0 to 51.25 (SD 19.04) at week 12. As per the previous two outcomes the control group registered worsening of their score by 5% over 12 week (45.94 [SD 21.36] at week 0 to 48.44 [SD 22.70] at week 12), P= 0.032, ES d= 0.11. These changes are graphically represented in Figure 4.





• Page 3 of 5 •

Muscle strength

Hip Abductors: Muscle strength of the hip abductors for the control group improved slightly on both sides, but did not reach statistical significance (Right hip mean baseline value= 5.98 [SD 2.67] to 6.20 [SD 2.74] after 12 weeks, P=0.606, ES d=0.08; Left hip mean baseline value= 5.85 [SD 3.11] to 6.23 [SD 3.30], P=0.210, ES d=0.12). The intervention group had significant improvements in muscle strength of the left side and despite not statistically significant, a marked improvement on the right (Right hip mean baseline value= 7.23 [SD 3.64] to 7.71 [SD 3.04] after 12 weeks, P=0.381, ES d= 0.14; Left hip mean baseline value= 6.02 [SD 2.99] to 7.33 [SD 2.94] after 12 weeks, P=0.014, ES d= 0.44).

Hip Adductors: A statistically signifcant decline in hip adductor muscle strength on both the right and left side (Right hip mean baseline value= 5.22 [SD 1.99] to 4.41 [SD 0.93], P=0.038, ES d= 0.53; Left hip mean baseline value= 4.39 [SD 1.67] to 3.74 [SD 1.21] after 12 weeks, P=0.014, ES d= 0.45) was noted for the control group. Statistically significant improvements were then obtained for participants in the intervention group (Right hip mean baseline value= 4.31 [SD 1.84] to 5.62 [SD 2.57], P=0.024, ES d= 0.59; Left hip mean baseline value= 4.12 [SD 2.83] to 5.73 [SD 2.83] after 12 weeks, P=0.010, ES d= 0.57).

Hip Extension: Participants in the control group registered a non statistically significant decline in muscle strength of the hip extenstors (Right hip mean baseline value= 4.09 [SD 2.84] to 3.78 [SD 2.45], P=0.376, ES d= 0.12; Left hip mean baseline value= 4.54 [SD 2.69] to 4.17 [SD 2.39], P=0.323, ES d= 0.46). A highly statistically significant increase in hip extensor muscle strength was registered for the intervention group (Right hip mean baseline value= 3.27 [SD 2.10] to 5.91 [SD 2.41], P<0.001, ES d= 1.17; Left hip mean baseline value= 3.69 [SD 2.54] to 5.88 [SD 2.84], P=0.003, ES d= 0.81).

Hip Lateral Rotators: Muscle strength for the hip lateral rotators in participants of the control group resulted in a deterioration, with the right side having a statistically significant decline in muscle strength as opposed to the left side (Right hip mean baseline value= 5.01 [SD 1.93] to 4.46 [SD 1.52] after 12 weeks, P=0.035, ES d= 0.32; Left hip mean baseline value= 4.76 [SD 2.05] to 4.32 [SD 1.53] after 12 weeks, P=0.233, ES d= 0.24). The intervention group, registered a statistically significant improvement recorded on the right side but not on the left despite an improvement in strength too (Right hip mean baseline value= 4.84 [SD 1.42] to 6.01 [SD 2.37] after 12 weeks, P=0.014, ES d= 0.6; Left hip mean baseline value= 4.91 [SD 1.36] to 5.71 [SD 2.06] after 12 weeks, P=0.107, ES d= 0.46).

Discussion

A 12 week exercise programme for patients with hip OA resulted in statistically significant improvements in most outcome measures with participants enrolled into the control group showing minimal improvements.

A statistically significant, 22% decline in pain scores for participants in the intervention group was noted (P=0.035, ES d=0.49) whereas participants enrolled into the control group had a slight increase in their pain levels by 0.5 (50% increase). 12weeks of exercises therefore resulted in an ideal period to lead to improvements in pain, to the contrary of what Uusi-Rasi et al [24] reported. These authors stated that 12 weeks might be a short time for effective progression in training, since this might lead to a rapid increase in exercise intensity

which could have adverse effects on pain levels, one which might dishearten participants to continue with treatment. Patients enrolled in the intervention group were being constantly monitored along the course for intensity progression which might have been one of the factors leading to such an improvement. Results obtained through this study are supported by Hernandez et al [25] who reported a significant reduction in pain behaviour in their participants, an intervention based on a three-times weekly sessions for 12 weeks, a protocol which was similar to what was carried out in this research study. A 12-week exercise intervention also matches recommendations issued by the American College of Sports Medicine (ACSM) pertaining to strength training. Hernandez et al (25) noted a marked reduction in the use of NSAIDs with these authors stating that despite this effect being a short-lasting one, the exercises might help in reducing or delaying the need of surgical intervention. Although not yet conclusive as to whether 12 or more supervised sessions are superior, it is reported that pain and physical function appear to improve further when this time frame is taken into consideration [26,27].

With a reduction in pain levels, the other outcomes also showed positive outcomes. A 12 week rehabilitation programme resulted in a relatively marked improvement in lower limb muscle strength for participants in the intervention group, but not for the control group. These benefits can be linked to the exercise progamme composition and also the duration of the programme. The revised 2017 clinical practice guidelines of the American Physical Therapy Association, highly consider with a grade A, the use of flexibility, strengthening and endurance exercises. Since muscle strength and pain levels improved, it is expected that the impact of OA on the individual will be ameliorated and also functionality. This could be seen from the 10.6% decline in WOMAC scores for the intervention group, ES d=0.32. The control group, on the otherhand had an increase of 5.4% in the WOMAC score.

A change in WOMAC score of 7.9 has been reported to be a minimal clinical important improvement in patients with hip OA as reported by Tubach [28] a value which was obtained in participants of the intervention group. The degree of short-term improvements in ADLs, physical performance and quality of life has been related by Hoglund et al [29] to the volume of supervised exercise received by the participants. Patients in the intervention group also reported a gain of 43m in the 6MWD, having a small magnitude ES of d=0.34. The control group on the other hand scored 1m less in distance from baseline in their 6-minute walking distance. Similar findings were reported by Widmer et al [30].

This study had some limitations one of which was the lack of proper monitoring of the HEP compliance using a diary/telephone communication, aspects which would have helped assess continuity of the exercises prescribed. The sample size for this study was also a small one and large sample groups will provide a better outline to outcomes from such an intervention.

Conclusion

This 12-week exercise intervention on a group of participants suffering from symptoms related to hip OA, resulted in clinical and statistically significant improvement in pain levels, functionality and lower limb strength. Such benefits follow the 2017 clinical practice guidelines for the use of flexibility, strengthening and endurance exercises in the rehabilitation for patients with hip arthritis.

• Page 4 of 5 •

Conflicts of interest

The authors declare that there are no conflicts of interest.

Funding

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• Page 5 of 5 •



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