

1 **Abstract**

2 **Purpose:** This study compares lower limb muscle strength and endurance in adults with hip
3 osteoarthritis, to an age-matched control group. **Methods:** Thirteen adults with moderate-to-severe
4 hip osteoarthritis (as graded by the Oxford Hip Score) and fifteen older adults participated.
5 Maximal voluntary isometric contraction of the knee extensors, knee flexors and hip abductors
6 and isotonic endurance of the knee extensors were measured using a dynamometer. Function was
7 assessed using the 30-second chair stand test, the 40 metre fast-paced walk test and a stair
8 negotiation test. Data were compared between groups using *t*-tests. **Results:** Participants with hip
9 osteoarthritis demonstrated weakness in the affected limb when compared to the control limb
10 during knee flexion (34%, $p=0.004$) and hip abduction (46%, $p=0.001$). Weakness was also
11 observed in the contralateral knee flexors (31%, $p=0.01$). When compared to the control limb, the
12 knee extensors of the hip osteoarthritis group were exhausted prematurely in the affected (70%,
13 $p=0.001$) and contralateral limb (62%, $p=0.005$). The hip osteoarthritis group took twice as long
14 to stair climb ($p=0.002$), walked 40% slower, ($p<0.001$) and had a 35% lower sit-stand
15 performance ($p<0.001$). **Conclusions:** Moderate-to-severe hip osteoarthritis may be characterised
16 by bilateral deficits in lower-limb maximal strength, markedly lower knee extensor endurance and
17 impaired functional performance.

18 **Keywords:** hip osteoarthritis; muscular strength; muscular endurance; physiotherapy; exercise;
19 function

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24 **Introduction**

25 Osteoarthritis is a common musculoskeletal condition within older adults, with an estimated 33%
26 of people aged 45 years and over having sought treatment for the disease in the United Kingdom
27 (UK) [1]. In the UK, 10.9% of adults aged over 45 years have osteoarthritis of the hip [1], which
28 often leads to joint pain, stiffness, reduced range of motion, slower gait speed and muscle weakness
29 [2-6]. Exercise is an integral component of the non-pharmacological management of osteoarthritis,
30 with local muscle strengthening and aerobic exercise recommended irrespective of age,
31 comorbidity, pain severity or disability [7]. Likewise, when progression of the disease leads to
32 consideration for total hip replacement surgery, preoperative physiotherapy and exercise
33 programmes (namely ‘prehabilitation’) are proposed as a potential method to expedite recovery
34 time and improve overall extent of recovery [8-10].

35 To develop effective physiotherapy and exercise programmes in osteoarthritis, it is crucial to
36 understand the underlying muscular impairment, and its relationship with physical function and
37 disease progression [11]. Whilst several research efforts have addressed maximal muscular
38 strength deficits in the hip osteoarthritis population [3], local muscular endurance has not been
39 studied to the same extent. Nonetheless, the physiological stimuli directed to skeletal muscle as a
40 result of strength training and endurance training are divergent in nature [12]. Maximal strength
41 involves exerting a maximum amount of force for a short period of time whereas muscular
42 endurance is the ability of the muscle or muscle group to sustain repeated contractions against a
43 load for an extended period of time [13].

44 Physiological studies have shown within a single muscle, motor units can vary greatly in their
45 contractile speed, maximum force, and resistance to fatigue [14]. These varying properties are co-
46 varying, such that the slowest units tend to be fatigue resistance but weaker (type 1 fibres), and the

47 strongest fibres are fast but relatively sensitive to fatigue (type II fibres) [14]. Muscle fibres can
48 adapt to changing demands by changing size or fibre type composition, and these changes may be
49 partially responsible for impairments and disabilities observed in patients who are deconditioned
50 [15]. Different fibre types enable muscles to fulfill a variety of functional demands [16], and the
51 endurance ability of the quadriceps femoris has a significant role for functional capabilities during
52 activities of daily living, such as walking and climbing the stairs [17]. Nonetheless, in chronic
53 disease, such as osteoarthritis, muscle fatigue may occur prematurely and persist [18]. The aim of
54 this study is to compare lower limb maximal muscle strength and local muscular endurance in
55 adults with hip osteoarthritis, to an age-matched control group.

56 **Methods**

57 *Participants*

58 This is an observational case-control study recruiting two study groups: i) adults aged over 60
59 years with a clinical diagnosis of unilateral or bilateral hip osteoarthritis and ii) healthy adults aged
60 over 60 years (control group) between 12th November 2019 and 15th March 2020. The study was
61 closed prematurely due to the COVID-19 pandemic and the Government advice to close higher
62 education institutes. Participants were recruited from the local area through online advertisement
63 (Twitter, Facebook) and email recruitment sent to local organisations. Sixty years was chosen as
64 the minimum age for the control group as osteoarthritis of the hip increases between the ages of
65 45 and 75 [1], and the average age for total hip replacement surgery is 68.0 ± 11.4 years [19].
66 Participants were included in the hip osteoarthritis group if they had: i) received a clinical diagnosis
67 of hip osteoarthritis from their general practitioner, an orthopaedic specialist or a physiotherapist;
68 ii) presented with chronic joint pain for at least three months; iii) had an Oxford Hip score [20] of
69 less than 40; and iv) were not on the waiting list for total hip replacement surgery. Participants

70 were included in the control group if they: i) were over 60 years old, ii) had no hip or lower limb
71 pain, iii) had no significant musculoskeletal comorbidities and iv) had no neurological diseases.
72 Exclusion criterion for both groups included: i) neurological disease affecting walking ability; ii)
73 rheumatoid arthritis; iii) fitted with a pacemaker or other active medical implant; iv) not physically
74 able to complete the testing protocol or v) not able to provide informed consent. The experimental
75 protocol was approved by the institutional ethics committee on 5th September 2019. In keeping
76 with good practice, the ethical principles for medical research outlined in the Declaration of
77 Helsinki were followed [21]. The STROBE (Strengthening the Reporting of Observational Studies
78 in Epidemiology) statement for the reporting of cross-sectional studies was used to guide the
79 reporting of this study [22].

80 *Variables*

81 Participants were invited to attend a laboratory-based testing session. Age, weight, height, and
82 medical history were recorded from all participants. Affected side(s), duration of symptoms and
83 the use of analgesia for pain relief were recorded from the participants in the hip osteoarthritis
84 group. The subjective severity of hip pain when weight bearing was rated using the Numeric Pain
85 Rating (NPR) scale (range 0-10 with 0 depicting minimal pain and 10 representing unbearable
86 pain) and the severity of symptoms were quantified using the Oxford Hip Score [20]. Physical
87 activity levels were collected using the Physical Activity Scale for the Elderly (PASE)
88 questionnaire [23].

89 *Maximal voluntary isometric contraction of the lower limbs*

90 Maximal, voluntary, isometric contraction (MVIC) of the lower limbs was measured using a
91 multimodal dynamometer (Primus RS, Baltimore Therapeutic Equipment, Hanover, USA) in line

92 with an experimental methodology published by our institute [24]. Muscle contractile force
93 estimations were conducted on both legs of the participants, with one leg tested at a time.

94 Participants were asked to perform three repetitions of three second maximal contractions of knee
95 extension, knee flexion and hip abduction, to encompass the muscle groups often affected by hip
96 osteoarthritis and most relevant to functional capacity [3,4,25]. For knee flexion and extension,
97 participants were seated on a secure chair with an 110° angle between the seat and the back of the
98 chair and with their knee flexed at 90° [24]. Participants were secured with a Velcro strap
99 positioned around their hips to limit contralateral compensation. The pivotal point of the lever was
100 aligned with the rotation of the knee joint to maintain appropriate position during all testing. To
101 perform knee flexion, the centre of the dynamometer pad was placed on the posterior part of the
102 leg, 5cm, above the lateral malleolus. For knee extension, the dynamometer exercise head was
103 applied to the anterior tibia, 5cm above the lateral malleolus [26]. Whilst there is much debate on
104 whether hip abduction should be performed in standing, side lying or supine [27,28], standing has
105 been defined as the most physiological [29] and functional [30] position for hip abduction
106 assessment, as the majority of daily activities involves hip abduction performed in weight bearing
107 conditions [31]. Hence, hip abduction was measured in the standing position, with the centre of
108 the dynamometer pad located 5cm proximal to the lateral femoral condyle [31]. The participant
109 began in the neutral position and then progressed to 10° abduction of the hip joint. Both arms were
110 placed in a neutral position, and a chair was placed in front of the participant in case they required
111 stabilisation during the test. Force was automatically adjusted by the dynamometer to account for
112 the length of the dynamometer attachment and lower limb segments distal to the joint being tested
113 [24].

114 Before each maximal series, participants carried out a warm-up, followed by one trial submaximal
115 effort of isometric knee flexion and extension and hip abduction, to become familiar with the
116 contractions. The warm-up included five minutes of level walking in the lab at a self-selected and
117 comfortable maximal speed, followed by a muscle warm up of both legs against a moderate
118 resistance (approximately 25% of the participants' body weight), applied by the dynamometer.
119 Participants with osteoarthritis completed the tests on their contralateral side first for
120 familiarisation purposes. Participants were given consistent verbal encouragement during each
121 contraction for attainment of maximal performance [32] and a one-minute recovery period was
122 observed between each maximal effort [33]. If the coefficient of variation of the three tests was
123 greater than 12.5%, the test was repeated following a period of rest to improve reliability of the
124 study findings [34]. A mean value from the three efforts was recorded for MVIC, and later
125 normalised to body mass (kilograms (kg)) (Nm/kg), to account for the confounding influence of
126 body weight on dynamometric measurement [35].

127 *Isotonic muscular endurance of the lower limbs*

128 Following a five-minute rest period, dynamic lower limb endurance was measured on the
129 dynamometer by calculating total energy expenditure (in joules) during repetitions of knee
130 extension/flexion at a constant cadence under a resistance of 40% of MVIC. The knee extensor
131 muscle group was chosen due to the significant role of the quadriceps muscle endurance for
132 functional capabilities during activities of daily living [17], and an isotonic movement was chosen
133 to best replicate functional activity. Participants were instructed to contract against the resistance
134 throughout the desired arc of motion, and to complete as many repetitions as possible, at the set
135 speed, calculated by the dynamometer as distance divided by time. Performance of each repetition
136 was monitored through the dynamometer power output. A successful repetition consisted of

137 completing the entire arc of motion within 2.5 seconds. When this criterion was not achieved, the
138 dynamometer power output would drop. A drop of $\leq 75\%$ power output (as compared with the first
139 repetition) was considered a failed repetition [36]. The test was ended after two successive failed
140 repetitions or the participant reported exhaustion and asked to stop. Participants were seated in
141 accordance to the positioning described for the MVIC knee extension test and secured with a
142 Velcro strap. Endurance was tested on both legs, one at a time, with the contralateral side tested
143 first in the osteoarthritis group, for familiarisation purposes.

144 *Functional assessment*

145 Functional performance was assessed through the 30-second chair stand test, the 40 metre (m) fast-
146 paced walk test and a stair negotiation test, as recommended as the minimal core set of
147 performance-based tests of physical function for older individuals (>40 years) diagnosed with hip
148 osteoarthritis [37]. The 30-second chair stand test measured the number of times the participant
149 could rise fully from chair and return to the seated position in 30 seconds. The participants' arms
150 were crossed at the wrist and held close to the chest to avoid upper body compensation. Walk
151 speed was measured as the time taken (in seconds) to complete a 10-metre walkway four times as
152 quickly as possible but at a safe pace. Stair negotiation was measured as the time taken in seconds
153 to safely ascend and descend eleven stairs (including turning around at the top) with a 20cm (8inch)
154 step height, at a self-selected pace. A handrail was provided but not used unless necessary for
155 safety.

156 *Sample size and statistical methods*

157 Since no preliminary data were available, sample size estimates were determined using data from
158 the knee extension MVIC scores for the first eight subjects tested in each group. Using the means

159 and standard deviation of the MVIC scores, our effect size was 0.96. Using a type 1 error protection
160 of 0.05, and a power of 0.80, we anticipated that 15 participants in each group to detect an outcome.
161 All data were analysed using IBM SPSS Statistics version 26 (SPSS Inc., Chicago, USA), with the
162 significance level set at $p < 0.05$. Both the right and left sides of the control group were tested to
163 observe any asymmetries, and then the right side was used as the comparison limb. The hip
164 osteoarthritis group were analysed per affected or contralateral side. For those with bilateral hip
165 osteoarthritis ($n = 2$), the affected or contralateral side were determined by the most and lesser
166 affected hip. Data normality was evaluated using a Shapiro-Wilk test. If both samples passed the
167 preliminary normality test, an independent samples or paired t test was conducted [39] to evaluate
168 differences between groups and between legs, respectively. Patient demographics were assessed
169 using an independent samples t test (age, BMI), a Mann Whitney U (PASE) or Fisher's exact test
170 (gender distribution). Mean (standard deviation) and median (interquartile range (IQR)) were used
171 to describe normally and non-normally distributed data, respectively [40]. Percentage change was
172 also used to describe the differences between the control and osteoarthritis group. Effect sizes for
173 differences in population means were computed using Cohen's d [41].

174 **Results**

175 *Participants*

176 Fifty-eight individuals volunteered to take part in the study (figure 1). During the initial telephone
177 consultation, 16 volunteers did not meet the inclusion criteria due to: musculoskeletal comorbidity
178 ($n = 6$); prior joint replacement ($n = 5$); hip pain but no clinical diagnosis of osteoarthritis ($n = 2$);
179 cardiovascular comorbidity ($n = 1$), fitted with a pacemaker ($n = 1$); and listed for total hip
180 replacement surgery ($n = 1$), and were excluded from the study. Six participants declined

181 participation due to travel or time commitments. A total of 36 were invited to attend the testing
182 session. Two participants in the control group were excluded during the eligibility assessment due
183 to knee pathology not previously disclosed. A further six participants were unable to attend the
184 testing session due to the COVID-19 pandemic and the Government advice to close higher
185 education institutes. Hence, the study was prematurely closed on 15th March 2020. This analysis
186 includes 28 participants who were recruited prior to the pandemic (hip osteoarthritis, $n = 13$;
187 control group, $n = 15$). Although we anticipated that 15 participants in each group to detect an
188 outcome, a post-hoc power analysis suggested that it was only the knee extension MVIC measure
189 that was underpowered.

190 There were no differences between groups in terms of age ($p = 0.39$) or gender distribution ($p =$
191 1.00). The osteoarthritis group were less active than the control group (median: 121.63 (IQR 70.80
192 to 175.13), versus 172.16 (IQR 95.07 to 209.71)), although this difference was not significant (p
193 $= 0.11$). The hip osteoarthritis group had a significantly higher BMI than the control group ($p =$
194 0.03). Participants with hip osteoarthritis group had a mean Oxford Hip Score of 28 ± 8 (range: 18
195 $- 39$), suggesting moderate-to-severe hip osteoarthritis [20]. As graded by the Oxford Hip Score,
196 one participant had severe symptoms (Oxford Hip Score of 18), six participants had moderate-to-
197 severe symptoms (Oxford Hip Score of 20-29), and six participants had mild-to-moderate
198 symptoms (Oxford Hip Score of 30-39). The mean duration of symptoms was 4 ± 3 years (range:
199 6 months $- 10$ years) and mean VAS pain on weight bearing was 5.31 ± 1.49 (range 3 $- 8$) (table
200 1). Eleven participants had unilateral hip osteoarthritis, and two bilateral. The two participants with
201 bilateral osteoarthritis did not produce any anomalies in the results. Six participants were not taking
202 any analgesics, four were taking paracetamol or ibuprofen when required, one was taking codeine

203 and paracetamol, one was taking the maximum dose of paracetamol, and one participant was taking
204 diahydrocodine in addition to cod liver oil.

205 [Insert Table 1]

206 [Insert Figure 1]

207 ***Variables***

208 *Maximal voluntary isometric contraction of the lower limbs*

209 One participant was not able to complete hip abduction MVIC on their contralateral side due to
210 pain in their affected hip. There were no differences in maximal strength of the left and right legs
211 of the control group (knee extension, $p = 0.67$; knee flexion, $p = 0.65$; hip abduction, $p = 0.06$)
212 whereas some asymmetries were observed in the hip osteoarthritis group. The affected leg of the
213 osteoarthritis group was 10% and 35% weaker when compared to the contralateral side in the knee
214 extensors ($p < 0.001$) and hip abductors ($p = 0.003$), respectively. No asymmetry was observed in
215 the knee flexors (table 2, figure 1).

216 In the osteoarthritis group, the knee extensors (quadriceps femoris) of the affected leg
217 demonstrated weakness when compared to the control group (22%), although this was not
218 significant ($p = 0.07$), perhaps due to this variable being underpowered. Similarly, the contralateral
219 leg was 14% weaker during MVIC knee extension when compared to the control group, but also
220 did not reach significance ($p = 0.23$). The knee flexors (hamstrings, gracilis, sartorius,
221 gastrocnemius, plantaris and popliteus) of the affected leg were 34% weaker than the control group
222 ($p = 0.004$), and the contralateral side demonstrated a 31% weakness in MVIC ($p = 0.01$), (effect
223 sizes 1.20 and 1.07, respectively). The hip abductors (gluteus medius, gluteus minimus and tensor
224 fasciae latae) demonstrated the most substantial weakness when compared to the control group,

225 with a 46% strength deficit in the affected side ($p = 0.001$, effect size 1.41), whilst the contralateral
226 side was only 18% weaker ($p = 0.22$, effect size 0.99).

227 [Insert Figure 2]

228 *Isotonic muscular endurance of the lower limbs*

229 No asymmetries were observed between the isotonic endurance of the knee extensors of the
230 affected and contralateral legs of the participants with hip osteoarthritis ($p = 0.26$) or left and right
231 legs of the control group ($p = 0.12$). In the osteoarthritis group, isotonic muscular endurance of the
232 knee extensors in the affected leg was 70% lower than the right leg of the control group ($p = 0.001$).
233 Likewise, the knee extensors of the contralateral leg were exhausted prematurely when compared
234 to the right leg of the control group (62%) ($p = 0.005$) (figure 3). Both comparisons yielded large
235 effect sizes (1.41 and 1.17, respectively). One participant in the hip osteoarthritis group reported
236 exhaustion before a drop of $\leq 75\%$ power output was observed on the dynamometer, and one
237 participant reported pain and asked to stop.

238 [Insert Figure 3]

239 [Insert Table 2]

240 *Functional assessment*

241 Participants with hip osteoarthritis score significantly worse than the control group in all three
242 functional tests, yielding very large effect sizes (table 3).

243 [Insert Table 3]

244

245

246 **Discussion**

247 This study compared unilateral maximal isometric strength and muscular endurance of the affected
248 and contralateral lower limbs in individuals with symptomatic hip osteoarthritis (graded moderate-
249 to-severe by the Oxford Hip Score) to a healthy, age-matched control group. Individuals with hip
250 osteoarthritis had weaker knee flexors and hip abductors than the control group in terms of MVIC
251 in both the affected and contralateral sides. In addition, participants with hip osteoarthritis
252 demonstrated lower isotonic muscular endurance of the knee extensors in both the affected and
253 contralateral limbs. The knee extensors also demonstrated weakness in terms of MVIC, although
254 this was not significant, perhaps as this outcome measure was underpowered. The deficits in
255 strength and endurance were consistent with the results of the functional assessment; whereby
256 participants with hip osteoarthritis performed significantly worse than their healthy counterparts
257 in all three tests. Asymmetries were observed in the maximal isometric strength of the knee
258 extensor and hip abductors in the affected and contralateral legs of the hip osteoarthritis group.

259 Comparison of the present study to the evidence-base is difficult due to differences in participant
260 characteristics, disease progression, maximal strength testing methodologies and a paucity of
261 evidence in the area of muscular endurance and hip osteoarthritis. In a review published in 2013
262 [3], thirteen studies were found to evaluate muscle strength, size, and/or inhibition in participants
263 with hip osteoarthritis [42-54]. Individuals with hip osteoarthritis were found to exhibit generalised
264 muscle weakness of the affected limb, which was underpinned by a combination of muscle
265 atrophy, reduced muscle density and muscle inhibition relative to the contralateral leg and control
266 group [3]. The greatest reductions in strength of the affected leg compared with the contralateral
267 leg were found for the hip and knee extensors and flexors, followed by the hip abductors and
268 adductors [3]. However, all but two articles [43,44] recruited participants from the waiting list for

269 a total hip replacement and included patients with advanced hip osteoarthritis. A more recent meta-
270 analysis examined only the hip abductor muscles, and found weakness in the affected leg when
271 compared to a control group and the unaffected limb across eight studies [55].

272 Few studies have investigated strength deficits at earlier stages of the disease. In the study by
273 Zacharias et al., maximal hip abductor strength was 40% lower in the affected leg of individuals
274 with moderate-to-severe hip osteoarthritis, when compared to a control group [25], similar to the
275 46% deficits observed in the present study. In a study by Loureiro et al., mild-to-moderate hip
276 osteoarthritis was characterised by 22% less knee extensor strength [2], and Rydevik et al. reported
277 15% less strength in patients with mild-to-moderate pain, when compared to controls [56],
278 comparable to the 22% weakness observed here. No evidence of between-limb asymmetries in
279 muscle strength or volume were found in individuals with mild-to-moderate hip osteoarthritis [2],
280 whereas in the present study, asymmetries were observed in the hip osteoarthritis group. In line
281 with previous investigations, these findings suggest strength asymmetries are a characteristic of
282 advanced hip osteoarthritis [3].

283 The main and novel finding of this study is that endurance of the knee extensors was markedly
284 lower in both the affected and contralateral sides of the hip osteoarthritis group when compared to
285 the left and right sides of the control group. These findings are perhaps not surprising, given that
286 muscle atrophy in osteoarthritis is homogeneous among both fibre types [57], and the relationship
287 between maximal isometric strength and relative muscular endurance [58,59]. These findings may
288 also be explained by changes in muscle fibre composition, whereby muscle fibres adapt to
289 changing demands [15]. For example, muscle disuse, prominent in the osteoarthritis population,
290 can lead to slow switch muscle fibres changing to a fast-twitch, fatigable fibre type [15].
291 Nonetheless, knee extensor endurance has important clinical implications due to the significant

292 role of the quadriceps femoris endurance for functional capabilities during activities of daily living,
293 such as walking, rising from a chair, and climbing stairs [17]. Findings from the functional
294 assessment further highlight the need to train endurance in the lower limbs of individuals with hip
295 osteoarthritis, given that it took this group twice as long to complete the stair negotiation test, 40%
296 longer to complete the 40m walk test and they had a 35% lower sit-stand performance. These
297 findings may be a result of deconditioning of the participants with osteoarthritis, secondary to their
298 pain. Activity levels in the osteoarthritis group were lower than in the control group, and inactivity
299 can lead to deconditioning of the muscular, cardiovascular, and respiratory systems [60].
300 Furthermore, as both the affected and contralateral limbs fatigued sooner than the control limb,
301 deconditioning combined with pain are likely to be causing factors.

302 Endurance training induces central and peripheral adaptations that improve cardiovascular
303 function and the capacity of skeletal muscles to generate energy through oxidative metabolism
304 [61]. In both hip and knee osteoarthritis, knee extensor exercises are commonly prescribed,
305 however not always with the training principles required to promote endurance benefits [62].
306 Training with low repetitions and high resistance favours adaptations for strength, power and
307 hypertrophy, whereas training with high repetitions and low resistance increases muscular
308 endurance and appears more suitable for submaximal, prolonged contractions [63]. Given the
309 concern that high-intensity or high-load strength training may increase pain and joint stress for
310 those with osteoarthritis [64], in addition to the function and endurance deficits observed in the
311 present study, endurance training may be the most suitable training modality in the hip
312 osteoarthritis population. For example, research has shown benefits of indoor cycling classes
313 [65] and circuit-based weight training for adults with hip osteoarthritis [66]. Clinically, our

314 findings are important to inform specific exercise prescription in physiotherapy and exercise
315 programmes for the hip osteoarthritis population.

316 A clear limitation of this study is the failure to meet the sample size estimates calculated due to a
317 global pandemic and the premature completion of data collection. The ongoing pandemic and
318 closure of higher education institutes prevented us from re-opening the study and recruiting any
319 additional participants. However, a post-hoc power analysis suggested that it was only knee
320 extension MVIC that was underpowered. More females were recruited to the study than males
321 which may be a source of experimental bias. However, as approximately 60% of the osteoarthritic
322 population are female, and 40% male [1], the participants in this study offer a reliable
323 representation of the wider population. Similarly, whilst the participants with hip osteoarthritis had
324 significantly higher BMI than the control group, obesity is widely acknowledged as a risk factor
325 for both the incidence and progression of osteoarthritis [67]. In addition, obesity can be a
326 consequence of osteoarthritis due to reduced physical activity due to joint pain, and hence, the
327 sample may offer a true representation of the population. Nonetheless, strength measurements
328 were normalised to body mass, to account for the confounding influence of body weight on
329 dynamometric measurement. Finally, the cross-sectional design of this study does not allow us to
330 evaluate whether muscle weakness is a cause or consequence of hip osteoarthritis.

331 **Conclusion**

332 In addition to bilateral deficits in maximal strength of the hip and knee muscles, moderate-to-
333 severe hip osteoarthritis may be characterised by markedly lower muscular endurance of the knee
334 extensors and impaired functional performance. The endurance capacity of the knee extensors can

335 play an important role in daily function, and thus it is important to consider endurance training
336 principles when prescribing exercise in hip osteoarthritis.

337

338 **Declaration of Interest Statement**

339 The authors report no conflicts of interest.

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504 **Figure captions**

505 Figure 1. Participant recruitment through study

506 Figure 2. Maximal voluntary isometric contraction (Nm/kg) and endurance (joules) in adults with
507 hip osteoarthritis compared to controls (mean \pm standard error)

508 **indicates significance when compared to the mean of the affected limb ($p < 0.05$)*

509 Figure 3. Knee extensor isotonic endurance, expressed as total work (joules) before two
510 consecutive contractions power were $\leq 75\%$ of the first repetition, in adults with hip osteoarthritis
511 compared to healthy controls (mean \pm standard error)

512 **indicates significance when compared to the mean of the affected limb ($p < 0.05$)*

Tables

Characteristic	Unilateral hip OA <i>n</i> = 11	Bilateral hip OA <i>n</i> = 2	All hip OA <i>n</i> = 13	Control group <i>n</i> = 15
Age (years)	75 ± 7.69	72 ± 4.95	75 ± 7.30	72 ± 6.42
Males, n (%)	4 (36%)	1 (50%)	5 (38%)	5 (33%)
Height (m)	1.68 ± 0.08	1.70 ± 9.90	1.68 ± 0.08	1.68 ± 0.12
Weight (kg)	83.0 ± 18.29	91.00 ± 4.24	84.23 ± 17.01	71.85 ± 14.89
BMI (kg/m ²)	29 ± 6	32 ± 2	30 ± 6	25 ± 4
Oxford Hip Score	27 ± 7	34 ± 5	28 ± 7	N/A
Pain (VAS)	5.79 ± 1.62	5.5 ± 0.71	5.31 ± 1.49	N/A
Duration of symptoms (years)	3.68 ± 2.82	6.0 ± 5.66	4.04 ± 3.17	N/A

Table 1. Sample characteristics

Measure	Affected limb Mean ± SD	Contralateral limb Mean ± SD	Difference between affected limb and contralateral limb (95% CI)	<i>p</i>	Effect size	Control limb (right) Mean ± SD	Difference between affected limb and control limb (95% CI)	<i>p</i>	Effect size	Difference between contralateral limb and control limb (95% CI)	<i>p</i>	Effect size
MVIC knee extension	2.56 ± 1.18	2.83 ± 1.13	0.27 (0.16, 0.37)	<0.001*	0.23	3.28 ± 0.72	0.71 (-0.07, 1.50)	0.07	0.74	0.45 (-0.31, 1.21)	0.23	0.47
MVIC knee flexion	1.14 ± 0.43	1.19 ± 0.46	0.05 (-0.09, 0.19)	0.49	0.11	1.73 ± 0.55	0.59 (0.20, 0.98)	0.004*	1.20	0.54 (0.14, 0.94)	0.01*	1.07
MVIC hip abduction	0.30 ± 0.14	0.46 ± 0.18	0.16 (0.07, 0.25)	0.003*	0.99	0.56 ± 0.22	0.27 (0.12, 0.41)	0.001*	1.41	0.10 (-0.06, 0.27)	0.22	0.50
Isotonic knee extensor endurance	659 ± 402	829 ± 721	170 (-140, 480)	0.26	0.29	2204 ± 1494	1544 (695, 2394)	0.001*	1.41	1375 (471, 2278)	0.005*	1.17

Table 2. Maximal voluntary isometric contraction (Nm/kg) and endurance (joules) in adults with hip osteoarthritis compared to controls

**indicates significant difference when compared to osteoarthritis (affected) leg (p < 0.05)*

Measure	All hip OA <i>n</i> = 13	Control group <i>n</i> = 15	Difference in means (95% CI)	<i>p</i>	Effect size
30s chair stand (n)	7.92 ± 1.44	12.13 ± 2.77	4.21 (2.45, 5.97)	< 0.001*	1.91
40m fast-paced walk (s)	45.92 ± 9.85	32.80 ± 4.86	13.12 (7.22, 19.03)	< 0.001*	1.69
Stair negotiation (s)	23.77 ± 11.04	11.80 ± 2.46	11.97 (5.22, 18.72)	0.002*	1.50

Table 3. Functional assessment scores for adults with hip osteoarthritis, compared to healthy

controls (mean ± SD)

**indicates significant difference between study groups ($p < 0.05$)*