

Effects of dry needling in gluteus medius muscle in individuals with chronic ankle instability: a randomized single-blinded controlled clinical trial

Type

Research paper

Keywords

pain, dynamic balance, dry needling, ankle instability

Abstract

Introduction

Background: Lateral ankle sprain (LAS) is the most common musculoskeletal injuries in sport and general population. The goal of the present study was to observe the effectiveness of dry needling (DN) in gluteus medius muscle in patients with chronic ankle instability (CAI).

Material and methods

A two-arm (1:1), single-blinded (participants), randomized clinical trial was performed in 40 subjects with chronic ankle instability and were divided in two groups: intervention group who received one session of dry needling in the most hyperalgesic gluteus medius myofascial trigger point (MTrP), (n = 20) and control group (n = 20). Dynamic balance, pain intensity, pain pressure threshold (PPT) and ankle dorsiflexion range of motion (ROM) were assessed at baseline, post-intervention and a 1-week follow up.

Results

The experimental group reported significant differences with respect to the control group for the anterior and medial dynamic balance ($p = .001$), PPT -ATL ($p = .002$) and ankle dorsiflexion ROM ($p = .001$).

Conclusions

The findings of the present study suggested that the DN in the most hyperalgesic MTrP of the Gmed muscle may increase the anterior and medial dynamic balance, ankle ROM and PPT-ATL at short-term in individuals with CAI. Pain intensity benefits were reported in both groups. Future studies should consider DN as a possible intervention in conjunction with a physical therapy program for individuals with CAI.

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2 **instability: a randomized single-blinded controlled clinical trial**

3
4 **Abstract**

5 Background: Lateral ankle sprain (LAS) is the most common musculoskeletal injuries in
6 sport and general population. The goal of the present study was to observe the
7 effectiveness of dry needling (DN) in gluteus medius muscle in patients with chronic
8 ankle instability (CAI). Methods: A two-arm (1:1), single-blinded (participants),
9 randomized clinical trial was performed in 40 subjects with CAI and were divided in two
10 groups: intervention group who received one session of DN in the most hyperalgesic
11 gluteus medius myofascial trigger point (MTrP), (n = 20) and control group (n = 20).
12 Dynamic balance, pain intensity, pain pressure threshold (PPT) in the anterior talofibular
13 ligament (ATL) and ankle dorsiflexion range of motion (ROM) were assessed at baseline,
14 post-intervention and a 1-week follow up. Results: The experimental group reported
15 significant differences with respect to the control group for the anterior and medial
16 dynamic balance (p = .001), PPT -ATL (p = .002) and ankle dorsiflexion ROM (p = .001).
17 Conclusion: The findings of the present study suggested that the DN in the most
18 hyperalgesic MTrP of the Gmed muscle may increase the anterior and medial dynamic
19 balance, ankle ROM and PPT-ATL at short-term in individuals with CAI. Pain intensity
20 benefits were reported in both groups. Future studies should consider DN as a possible
21 intervention in conjunction with a physical therapy program for individuals with CAI.

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35 Introduction

36 Lateral ankle sprain (LAS) is the most common musculoskeletal injuries in sport and
37 general population.(1) In addition, the societal cost as high as \$6.2 billion annually.(2)
38 Patients who suffer a LAS episode commonly report a decreasing in quality of life, foot
39 and ankle osteoarthritis, a lack of lower limb functionality and thus, a decreasing in sports
40 performance.(2) A 70% of those individuals may develop chronic ankle instability (CAI),
41 a disorder featured by subjective instability, recurrent episodes of “giving away” and
42 sprains, loss of function, postural control, reflex function, mechanical and sensorimotor
43 disturbances.(3) Moreover, Terrier et al.(4) reported that an altered proprioception,
44 balance and motor control can be reported in patients with CAI. Several authors described
45 that patients with CAI have been an altered gait kinematics, disturbances in ligament
46 laxity, range of motion (ROM) alterations and synovial changes.(5)(6)(7) The summation
47 of this symptoms may predispose an increase in the risk of recurrent LAS associated to a
48 laterally deviated center of pressure and disturbances in balance during the gait
49 movement.(8)(9)

50 It has been reported that patients with CAI showed an increase of peroneus longus (PL)
51 activity prior to the initial ground contact.(10) In addition, in an unilateral drop jump
52 patients with CAI reported a decreasing of the PL muscle activation compared with
53 healthy individuals assessed by electromyography.(11) Hale et al.(12) argued that the
54 existence of disturbances in both peripheral and centrally mechanisms could be developed
55 after injury processes, which produces alterations in postural control after rehabilitation
56 programs. Moreover, authors suggest that a balance program training on lower extremity
57 could have benefits in the foot and ankle function and balance. Despite of most of the
58 research about muscular motor control and patterns have been carried out in proximal
59 ankle joints in subjects with CAI (e.g. decreasing muscle activation in muscles that
60 surrounding the ankle, knee and hip),(13) Webster and Gribble(14) showed a decreasing
61 of gluteus maximus (Gmax) muscle activity in patients with CAI during a single leg
62 rotational squat work.

63 Postural control plays an important role for the development of an adequate muscular
64 activation and a coordinated gait pattern. The Star Excursion Balance Test (SEBT) was
65 defined as a reliable and valid tool to assess the dynamic postural control between
66 individuals with and without ankle pathology.(15) In addition, patients with CAI showed
67 a postural control deficit compared with healthy individuals measured by SEBT.
68 Likewise, the Y-Balance test (YBT) was also defined a SEBT variation tool to assess the

69 dynamic balance of the lower limb with excellent reliability values (intra-rater ICC = 0.85
70 to 0.91 and inter-rater ICC = 0.91 to 0.99)(16) and it was employed in previous
71 studies.(17)

72 Ayotte et al.(18) reported that an appropriate Gmax and gluteus medius (Gmed) function
73 is necessary for maintaining postural stability during weight bearing activities. In this
74 line, Jaber et al. compared the postural control and electromyographic (EMG) activity
75 between individuals with and without CAI. The results showed that alterations in
76 proximal and distal muscle activity have negative effects in quality of movement and
77 postural control, which may lead long-term functional.(19) Thus, authors suggested that
78 interventions and training programs focused in hip and ankle muscles could have benefits
79 in the prevention and management in patients with CAI.

80 Myofascial trigger points (MTrPs) located at the Gmax and Gmed were described as a
81 result of muscle weakness, biomechanical disturbances of the lower limbs or lumbar
82 pathology.(20) MTrPs were defined as an “hyperirritable nodule in a taut band of skeletal
83 muscle which is palpable and tender during the physical evaluation”.(21) In addition,
84 MTrPs were related with overuse, motor control disturbances, injuries and pain. Two
85 **classifications** of MTrPs were defined: active MTrPs reproducing symptoms and referred
86 pain and, latent MTrPs had no symptoms but can be palpable in a clinical examination.
87 Regarding the diagnosis, active MTrPs performed a recognizable pain and local twitch
88 response to a needle penetration.(20)

89 Dunning et al.(22) argued that dry needling (DN) have benefits on the management of the
90 neuromusculoskeletal pain syndromes, such as the treatment of MTrPs. In addition,
91 Salom et al.(23) reported that DN is an effective approach for the treatment of sensory
92 and motor factors in MTrPs. Several authors showed the benefits of the DN in the pain
93 management, for example Hu et al.(24) conducted a meta-analysis in patients with low
94 back pain showing that DN was more effective than acupuncture for reducing pain
95 intensity. In the same line, a systematic review and a meta-analysis carried out by Gattie
96 et al.(25) reported that DN was **superior to sham** treatment for short- and mid-term follow
97 ups for musculoskeletal pain conditions. In addition, DN performed in the
98 sternocleidomastoid muscle have also shown an increase of the motor control of the
99 cervical muscles for a 1-month follow up in individuals with neck pain(26). Sánchez-
100 **Mila et al.(27) reported the benefits of DN added to a Bobath program in ROM, balance**
101 **and also reducing the spasticity in patients who had suffered a stroke.**

102 Several authors reported that an improvement of the motor control and function of hip
103 muscles, such as Gmax and Gmed could be benefits on the lower limb stability. In
104 addition, for the treatment or prevention in patients who develop CAI. Thus, the aim of
105 the present study was to determine the effect of a DN intervention in the most
106 hyperalgesic latent MTrPs of the Gmed, mainly related to ankle ROM, dynamic balance
107 and the pressure pain threshold (PPT) of the anterior talofibular ligament (ATL) and
108 MTrPs of the Gmed muscle compared with a sham intervention in individuals with CAI.
109 We hypothesized that individuals receiving DN would exhibit greater improvements in
110 ankle ROM, balance and PPT than those patients receiving a sham intervention.

111 **Methods**

112 ***Design***

113 A two-arm (1:1), single-blinded (participants), randomized clinical trial was performed
114 from September 2019 to January 2020 following the CONSolidated Standards of the
115 Reporting Trials criteria.(28)

116 ***Ethical considerations***

117 Previously, the Clinical Research Ethics Committee of Hospital de la Princesa (Madrid,
118 Spain) approved the study and was registered in Clinicaltrials.gov (NCT04108390). All
119 the participants signed the informed consent form before the beginning of the study.
120 Moreover, all the participants respected and took into account the Helsinki Declaration
121 and ethical standards for human experimentation.

122 ***Sample size calculation***

123 G*Power software was employed for the sample size calculation by the difference
124 between the intervention group and control group using the ATL-PPT (kg/cm²) variable
125 of a pilot study (n = 12) divided in two groups (mean ± SD), 6 subjects for the A group
126 (intervention) (4.42 ± 0.25) and 6 subjects for the B group (control) (4.22 ± 0.21). For the
127 sample size calculation, a power of 0.80, an α error of 0.05 and effect size of 0.86 with 1
128 tailed hypothesis were employed. In conclusion, a sample of 36 was calculated. However,
129 we could recruit sample of 40 individuals for this study.

130 ***Participants***

131 A total sample of 40 subjects with CAI based on the position statement of the
132 International Ankle Consortium(29) from a care center was recruited for the present study
133 and divided in two groups: intervention group (n = 20) and control group (n = 20). (Figure
134 1). Inclusion criteria were as follows: history of at least 2 recurrent ankle sprains with
135 inflammatory symptoms (e.g. pain, swelling), at least 2 episodes of “giving away” in the

136 6 months before study enrollment.(5) Subjects were excluded if they: had no history for
137 ankle sprain, history of vestibular disorders, lower limb surgeries in the previous 12
138 months (e.g. fractures or muscular tears), patients who received a physiotherapy, medical
139 o pharmacology treatment in the previous 3 months.(19)

140 ***Randomization and blinding***

141 Before the intervention, the random process was developed with the free software system
142 randomization.org with 1:1 allocation ratio and assigned the participants to the A group
143 (intervention) or B group (control). Therefore, patients for each group were do not known
144 which group they belong.

145 ***Interventions***

146 Participants included in this study received 1 intervention with 3 evaluations: pre-
147 intervention, post-intervention and at 1-week. We used the same needles (0.32x40mm)
148 for both interventions. In addition, the interventions were performed by the same therapist
149 with more than 10 years of experience in DN technique for the management of MTrPs.
150 Before the needle application, the area was disinfected with skin antiseptic. Immediately
151 after the intervention, the therapist applied pressure into the skin in order to prevent
152 excessive bleeding with a cotton bud.

153 ***Intervention group***

154 Only one session of DN to the intervention group on the ipsilateral Gmed muscle of the
155 CAI lower limb using Hong's "fast in" and "fast out" intervention with multiple rapid
156 needle insertion was applied following previous guidelines.(30)(31) Each participant
157 received the same DN treatment in the most hyperalgesic latent MTrP located on the
158 Gmed muscle.

159 ***Control group***

160 Following Pecos et al.(32) guidelines, the control group received the same protocol on
161 the ipsilateral Gmed muscle of the CAI lower limb, but the needle was inserted 1.5 cm
162 medially from the MTrP (outside the MTrP location).

163 ***Outcome measurements***

164 PPT was defined as the amount of pressure generated of the target point to be evaluated
165 until painful sensation appears.(33) For the present study, PPT was assessed from 0 to 10
166 kg/cm² with a mechanical algometer (FDK/FDN, Wagner Instruments, Greenwich, CT).
167 In addition, this algometer is reliable, sensitive and reproducible for latent MTrP
168 assessments.(34)(35)(36) The most hyperalgesic latent MTrP in the Gmed and the ATL

169 were assessed. The evaluation procedure was performed by the mean of three repeated
170 measurements with a 30-60s rest interval between evaluations.

171 The YBT consist of three lines attached to the floor in the anterior posteromedial and
172 posterolateral directions. Following the Pliski et al.(16) procedure, the posterior lines
173 were located 135 degrees from the anterior line with 45 degrees between the posterior
174 lines. Before the test, individuals viewed an instructional video about the procedure in
175 order to the familiarization process. Once the demonstration have been carried out, the
176 subjects practiced six trials on each leg of the three directions prior the formal
177 assessment.(16) The subjects were in standing barefoot at the center of the “Y” mark.

178 **Each participant should to maintain a single-leg stance of the target limb in order to reach**
179 **the maximum distance in anterior, posterolateral and posteromedial directions.** The
180 subject’s hands were placed on their hips and the stance heel should be remain in contact
181 with the ground. If during the assessment any criteria were violated, the trial should be
182 repeat. To calculate the normalized YBT values the individual’s leg length was measured
183 in supine position from the anterior superior iliac spine to the malleolus tibialis.(37) The
184 distance was quantified in centimeters. For the normalization the following formula was
185 applied dividing the mean reach distance by the individuals leg length and multiplying
186 by 100%.

187 Maximal ankle dorsiflexion ROM was evaluated using a standard manual goniometer and
188 were defined as the distance of the toe from the wall maintaining the contact between
189 wall and knee without lifting the heel. Thus, the individuals reach the final lunge position
190 at maximal dorsiflexion and the goniometer was aligned with the mobile branch at the
191 fibula and the stable branch aligned with the fifth metatarsal head.(38) For each
192 measurement, the evaluator passively moved the ankle from a neutral baseline position
193 to a dorsiflexion until a firm end-feel was bringing out.

194 All the outcome measurements were carried out by the same investigator (G.J.C).

195 ***Statistical analysis***

196 SPSS 23.0 software (IBM SPSS Statistics, Armonk-NY; IBM-Corp) was employed for
197 the statistical analysis. Kolmogorov-Smirnov test was used to assess normality data
198 distribution. Student *t* test was applied to test age, weight, height, BMI and Cumberland
199 Ankle Instability Tool (CAIT) differences between groups. In order to check the basal
200 values of the main variables the Student *t* test was also employed. To assess the effects

201 of intra-subjects (time) and inter-subject (treatment groups) values on the dependent
202 variables, a two-way analysis of variance (ANOVA) for repeated measures was
203 performed (considering the significance of the Greenhouse-Geisser correction when the
204 Mauchly test rejected the sphericity). The Tukey post-hoc test was employed for multiple
205 comparisons. Furthermore, the effect size was calculated by the Eta² coefficient. For non-
206 parametric data, Friedman test and Wilcoxon post-hoc analyses were employed for intra-
207 subject comparisons. Moreover, Mann-Whitney *U* test was employed for the comparisons
208 between groups. The level of significance was set at $P < 0.05$ with an α error of 0.05 (95%
209 confidence interval) and a desired power of 80% (β error of 0.2).

210 **Results**

211 Regarding the table 1, sociodemographic data did not show significant differences
212 ($P > .05$). Moreover, Student *t* test for the main variables reported significant differences
213 between groups for ATL-PPT ($P = 0.015$) and VAS ($P = 0.002$) baseline variables. The
214 rest of variables did not show significant differences ($P > .05$) between the intervention
215 and control group. Time interaction effects reported significant differences ($P > .05$) for
216 SEB anterior, SEB lateral, SEB medial, dorsiflexion ROM, ATL-PPT and Gmed PPT
217 variables. Significant differences were observed between groups for an increase of YBT
218 anterior ($P = .001$), YBT medial ($P = .001$), dorsiflexion ROM ($P = .001$) and ATL-PPT
219 ($P = .002$) in favor the intervention group with respect to the control group. In addition,
220 no significant differences were obtained for Gmed PPT ($P = .332$) variable. (Table 2) In
221 addition, Tukey post-hoc analysis reported significant differences ($P > .05$) between
222 baseline and post-intervention measurements for ATL-PPT, dorsiflexion ROM, YBT
223 anterior, YBT lateral and YBT medial variables for the experimental group. Significant
224 differences between baseline and 1-week for ATL-PPT, dorsiflexion ROM, YBT
225 anterior, YBT lateral and YBT medial variables in the experimental group. (Figure 2)

226 Regarding the pain intensity variable, Friedman test reported differences ($p = 0.001$) for
227 the experimental group and the control group ($p = 0.043$). Mann-Whitney U test reported
228 differences between groups ($p = 0.004$) but non-significant differences ($P > .05$) were
229 reported between groups at post-intervention and at 1-week follow up.

230 **Discussion**

231 To the authors' knowledge, this research study may be considered the first clinical trial
232 showing benefits in balance, ankle dorsiflexion ROM and ATL- PPT in individuals with
233 CAI who received a DN intervention in the most hyperalgesic MTrP at the Gmed muscle.
234 In addition, pain intensity benefits were found in both groups. These findings were in
235 accordance with previous studies who reported the benefits of DN in MTrP at the PPT
236 variable in individuals with temporomandibular disorders,(39) chronic shoulder pain,(40)
237 neck pain,(41) patients after knee arthroplasty,(42) or plantar heel pain.(43) Moreover,
238 the results of the present study reported significant differences for the increase of the
239 ATL-PPT immediately post-intervention and at 1-week follow up compared with the
240 control group. Regarding the pain intensity at the experimental group, benefits were
241 shown immediately after the DN application and at 1-week follow up as well as the
242 control group. In a systematic review and meta-analysis conducted by Liu et al.(44)
243 showed that DN intervention of MTrP may be recommended to relieve the pain intensity
244 at short- and medium-term in individuals with low back pain, but the effects with larger
245 follow-ups remains unclear. In addition, in a systematic review conducted by the same
246 authors 3 years before authors reported that DN was effective to relieve the pain intensity
247 in subjects with low back pain at post-intervention and reported that further research is
248 needed to improve the knowledge of the effectiveness with longer follow-ups in DN
249 interventions. Considering the positive effects of a DN approach in the PPT, prior studies
250 reported benefits at the increase in PPT levels in the masseter muscle in patients with

251 temporomandibular disorders,(39) cervical spine (C7) in subjects with chronic neck pain
252 with a DN treatment in the upper trapezius muscle,(45) or in individuals with unilateral
253 shoulder impingement syndrome with a DN in the upper trapezius muscle.(46) Our results
254 were related with prior studies that showed benefits in pain intensity and PPT in ATL at
255 short-term with the application of DN in MTrP. A possible explanation for the
256 effectiveness of the DN in pain variables could be related with the gait control mechanism
257 for the speedily penetration of the needle into a MTrP might stimulate the afferent sensory
258 fibers, which produce an inhibition in the dorsal horn of the spinal cord by blocking the
259 pain afferences developed in the MTrP nociceptor.(47)

260 Several authors reported the effectiveness of the DN technique (isolated or combined with
261 other therapies) in the ROM. For example, Onat et al.(48) found an increase of ROM after
262 the application of DN into the posterior paracervical muscle in individuals with
263 mechanical neck pain. Mendiguita-Gómez et al.(49) included a DN treatment for the
264 spastic shoulder muscles in individuals who had experienced a stroke and reported an
265 increase of the ROM. The present study showed the effectiveness of a DN approach
266 increasing the ankle dorsiflexion ROM, these findings could be related with the
267 improvement of the dynamic balance of the lower limb, which can help to increase the
268 ankle mobility. In addition, the results reported by several authors about the ROM after a
269 DN treatment did not match with what we found in our study. (50)(51) Therefore, further
270 research is needed in order to a better understanding of the possible relationship between
271 the motor control and ROM.

272 Regarding the dynamic balance, MacKinnon and Winter argued that the gait in the frontal
273 plane is regulated by the subtalar and hip joint.(52) Thus, a deficit in the motor control at
274 the muscles involved with the hip kinematics would influence the gait in individuals with
275 and without pathology. In addition, Jun Son et al.(5) suggested that the ankle joint corrects

276 small disturbances related with a deficient foot position, and the hip correct large
277 disturbances associated to a deficient foot position. Thus, the ankle and hip have to work
278 in synergy. When the foot adopts vulnerable positions or in an overcompensation ankle
279 kinematics (e.g. ankle sprains, forced inversion) the lower limb dynamic balance might
280 be corrected for an increase of muscle activity of the ankle evertors and hip abductors,
281 such the Gmed. Our results reported the effectiveness of a DN intervention in the most
282 hyperalgesic MTrP located in the Gmed muscle in individuals with CAI for the increase
283 in dynamic balance with respect to controls for the YBT anterior and medial variables. In
284 addition, anterior, medial and lateral YBT variables reported differences between
285 baseline and 1-week follow up in the experimental group. A possible explanation of these
286 results could be an increase in the motor control of the targeted MTrP at the Gmed muscle
287 and an improvement of the musculoskeletal pain conditions. Despite of in this study did
288 not performed a electromyography analysis, several authors related the effectiveness of
289 the MTrP DN with a muscle function improvement.(53)(54) The activation of 6 lower
290 extremity muscles was reduced in individuals with CAI – tibialis anterior (7%), peroneus
291 longus (4%) and vastus lateralis (4%)- and a decreasing – Gmed (4%) and Gmax (10%)
292 – suggested that patients with CAI could be affected the neural activation pathways.(5)

293 ***Clinical applications***

294 **The results of the present study do not provide a gold standard approach for CAI patients.**
295 **Thus, these findings aim to provide novel scientific evidence to the clinicians and**
296 **researchers suggesting that DN technique could be effective in conjunction with the**
297 **manual therapy or exercise programs in individuals with CAI. Nevertheless, future**
298 **studies should clarify the addition of DN to these physical therapy interventions.**

299 **Limitations**

300 Some limitations should be acknowledged in the present study. First, baseline significant
301 differences were found between groups at baseline for ATL-PPT and VAS, therefore the
302 results of the present study for these variables might be biased. Second, for the YBT only
303 one measure was taken for each variable. Third, an electromyographic evaluation did not
304 carried out in order to evaluate the muscle activity of the extrinsic foot and the Gmed
305 muscle. At last, the DN treatment was not observed in the medium or long term. Further
306 research is recommended in order to evaluate the muscle activity, temperature of the
307 MTrP or with a larger follow ups in patients with CAI.

308 **Conclusions**

309 The findings of the present study suggested that the DN application in the most
310 hyperalgesic MTrP of the Gmed muscle may increase the anterior and medial dynamic
311 balance, ankle dorsiflexion ROM and PPT-ATL at short-term in individuals with CAI.
312 Pain intensity benefits were reported in both groups. Future studies should consider DN
313 as a possible intervention in conjunction with a physical therapy program for individuals
314 with CAI. Further research is still needed to improve the knowledge about the pain
315 perception mechanisms developed in subjects with musculoskeletal disorders with
316 invasive physiotherapy approaches.

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510 **Figure legends.**

511 Figure 1. Flow chart diagram.

512 Figure 2. Comparison between intervention and control group measurements for YBT,
513 PPT and VAS variables. * Significant differences between groups (treatment x
514 time).

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Table 1. Sociodemographic data and CAIT scale of the sample

Data	Total sample (n = 40)	Intervention		P-value Cases vs Controls
		(n = 20)	Controls (n= 20)	
Age, y	32.50 ± 6.78	33.80 ± 6.77	31.20 ± 6.70	.230
Weight, kg	68.52 ± 12.02	70.60 ± 13.27	66.85 ± 10.65	.331
Height, m	1.69 ± 0.18	1.66 ± 0.25	1.73 ± 0.07	.220
BMI, kg/m ²	22.81 ± 1.35	24.61 ± 1.05	25.36 ± 1.47	.350
CAIT	22.30 ± 2.24	22.60 ± 2.03	22.00 ± 2.44	.405

Abbreviations: BMI, body mass index; CAIT, Cumberland ankle instability tool.

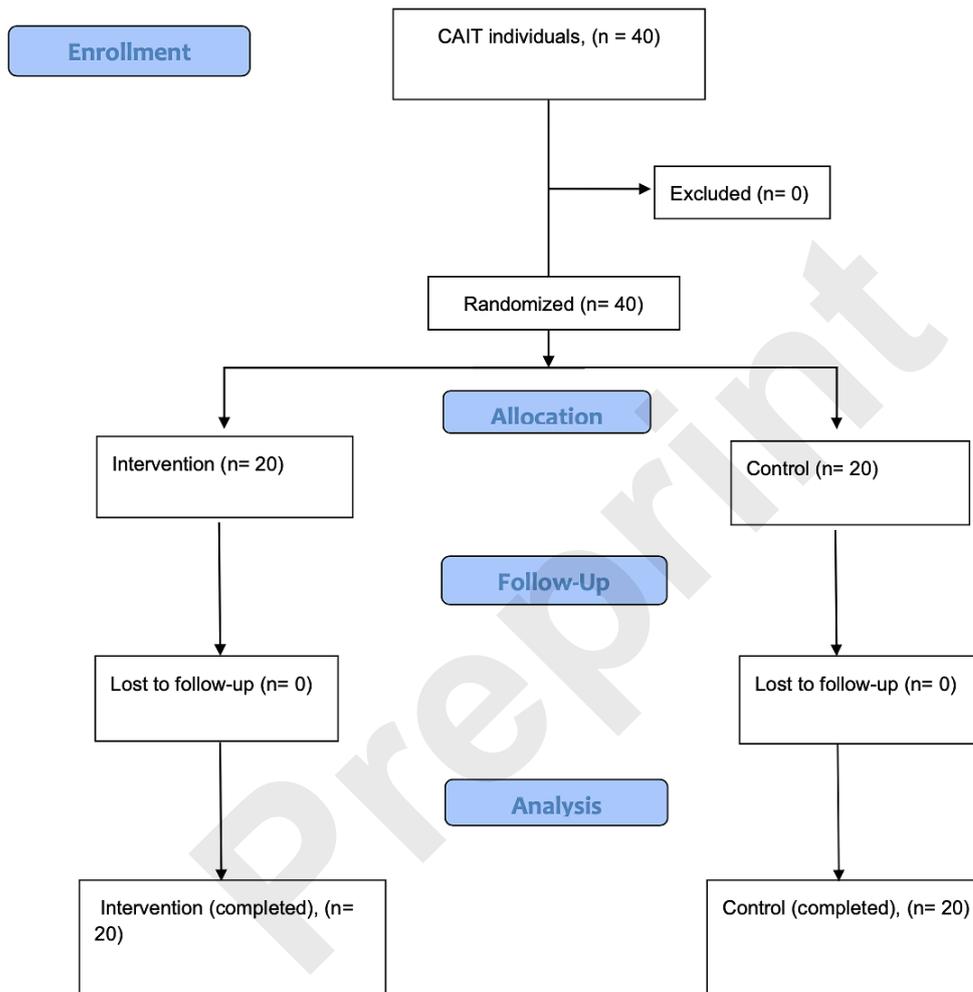
Table 2. SEB, ROM, VAS and PPT intrasubject effects.

Intrasubject Effects				
Measure	Intervention n=20	Control n=20	Time value F; P (Eta ²)	Treatment X Time F; P (Eta ²)
YBT anterior			F = 22.272; P = .001 (0.370)	F = 7.425; P = .001 (0.163)
Baseline	94.06 ± 24.78	97.56 ± 8.96		
Post-test	99.28 ± 24.56	98.52 ± 8.73		
1-week	100.68 ± 23.58	99.53 ± 8.95		
YBT medial			F = 28.391; P = .001 (0.428)	F = 15.075; P = .001 (0.284)
Baseline	131.06 ± 16.41	134.50 ± 15.92		
Post-test	138.80 ± 16.46	135.67 ± 15.30		
1-week	140.96 ± 15.43	136.07 ± 14.98		
YBT lateral			F = 9.650; P = .001 (0.203)	F = 3.245; P = .063 (0.079)
Baseline	120.83 ± 21.24	126.12 ± 17.21		
Post-test	126.08 ± 21.77	127.20 ± 16.96		
1-week	126.97 ± 19.26	127.96 ± 17.11		
1-week	77.0 ± 4.8	73.2 ± 6.2		
Dorsiflexion ROM			F = 8.436; P = .001 (0.182)	F = 7.728 P = .001 (0.169)
Baseline	149.15 ± 11.89	133.40 ± 13.77		
Post-test	152.85 ± 12.23	133.55 ± 13.53		
1-week	152.25 ± 11.03	133.35 ± 13.56		
ATL PPT			F = 40.613; P = .001 (0.517)	F = 9.372; P = .002 (0.198)
Baseline	3.21 ± 0.82	3.91 ± 0.89		
Post-test	3.92 ± 0.93	4.11 ± 0.98		
1-week	4.04 ± 0.96	4.23 ± 1.01		
Gmed PPT			F = 6.559; P = .009 (0.147)	F = 1.037; P = .332 (0.027)
Baseline	5.46 ± 2.16	5.65 ± 1.24		
Post-test	5.79 ± 2.33	5.77 ± 1.23		
1-week	5.91 ± 2.29	5.86 ± 1.18		

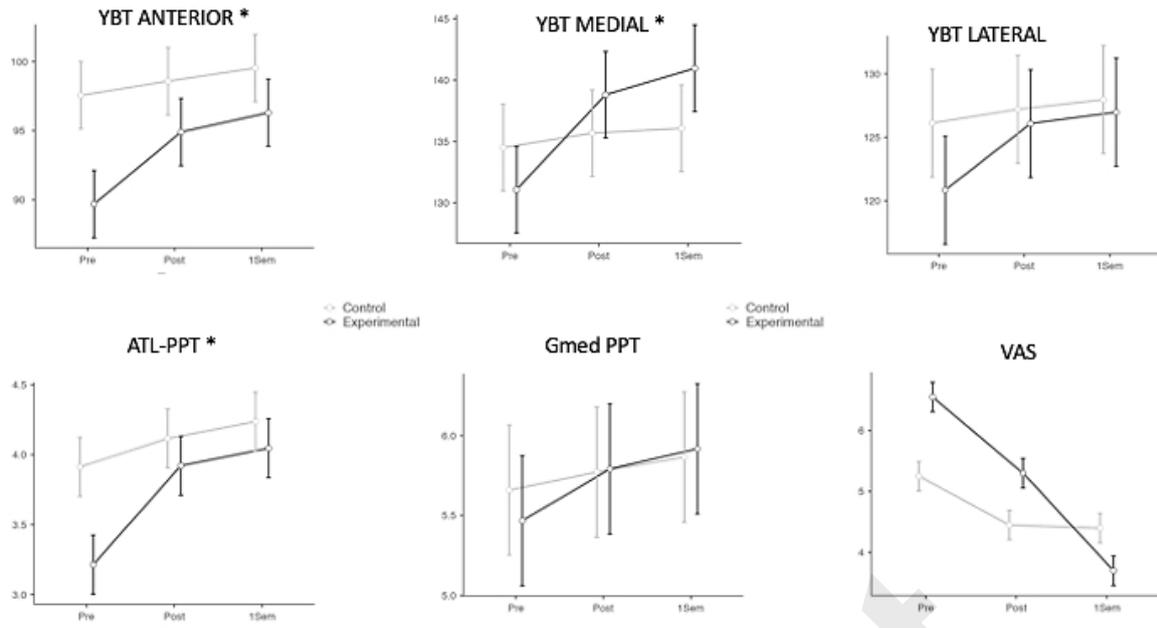
Values are mean ± SD unless otherwise indicated.

Abbreviation: ATL, anterior talofibular ligament; Gmed, gluteus medius; PPT, pain pressure threshold; ROM, range of motion; SEB, star excursion balance; YBT, Y-balance test.

CONSORT Flow Diagram



Flow chart diagram



Comparison between intervention and control group measurements for YBT, PPT and VAS variables. * Significant differences between groups (treatment x time).

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