Title: Gait Training Interventions for Individuals with Chronic Ankle Instability: A Systematic Review &

Meta-Analysis

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1 Gait Training Interventions for Individuals with Chronic Ankle Instability: A Systematic Review &

2 Meta-Analysis

3

4 Abstract

5 **Objective:** This review aimed to determine if gait training interventions influence lower

- 6 extremity biomechanics during walking in individuals with chronic ankle instability (CAI).
- 7
- 8 Methods: A literature search was conducted in PubMed, CINAHL, SPORTDiscus, and
- 9 MEDLINE to identify English-language studies from inception through September 2022.
- 10 Eligible studies included randomized control trials, repeated measures design, and descriptive
- 11 laboratory studies measuring the effects during or following a gait training intervention on
- 12 biomechanical outcomes (kinematics, kinetics, electromyography) during walking in individuals
- 13 with CAI. Gait training interventions were broadly categorized into devices (destabilization
- 14 devices, novel gait training device) and biofeedback (visual, auditory, and haptic delivery
- 15 modes). Meta-analyses were conducted when appropriate using random-effects to compare pre-
- 16 and post- gait training intervention mean differences and standard deviations.
- 17
- 18 **Results:** Thirteen studies were included. Meta-analyses were conducted for single session gait
- 19 training studies only. Eleven studies reported kinetic outcomes. Our meta-analyses showed
- 20 location of center of pressure (COP) was shifted medially from 0-90% (Effect Size [ES]
- range=0.35-0.82) of stance, contact time was decreased in medial forefoot (ES=0.43), peak
- 22 pressure was decreased for lateral midfoot (ES=1.18) and increased for hallux (ES=0.59),
- pressure time integral was decreased for lateral heel (ES=0.33) and lateral midfoot (ES=1.22)
- and increased for hallux (ES=0.63). Three studies reported kinematic outcomes. Seven studies
- reported electromyography outcomes. Our meta-analyses revealed increased activity following
- 26 initial contact (IC) for fibularis longus (ES=0.83).
- 27
- 28 Conclusions: Gait training protocols improved some lower extremity biomechanical outcomes 29 in individuals with CAI. Plantar pressure outcome measures seem to be most impacted by gait 30 training programs with improvements in decreasing lateral pressure associated with increased 31 risk for lateral ankle sprains. Gait training increased EMG activity post-IC for the fibularis 32 longus. Few studies have assessed the impact of multi-session gait training on biomechanical 33 outcome measures. Targeted gait training should be considered when treating patients with CAI.
- 33 34
- **Key Words:** Ankle sprain, biomechanics, biofeedback, rehabilitation, gait training device
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- Gait training improved lower extremity biomechanics associated with risk for lateral ankle sprains, including medial shifts in plantar pressure, decreased ankle inversion, and increased fibularis longus activity with medium to large effect sizes.
- Significant gait improvements were evident utilizing a variety of gait training devices and biofeedback.

47	•	There is limited evidence on kinematic outcomes of gait training interventions for
48		CAI.
49	•	Gait training would benefit from homogeneity between protocols and techniques
50		suitable for clinical implementation.
51		



52 Introduction

53 Lateral ankle sprains (LAS) are a prevalent musculoskeletal injury among the general population and physically active individuals.^{1,2} These injuries can be temporarily disabling, 54 hinder physical activity, and contribute to long-term ankle joint problems.^{2,3} Recurrent LAS rates 55 are high⁴ and in one prospective study, 40% of individuals who sustained their first ever LAS 56 developed a condition known as chronic ankle instability (CAI).⁵ CAI is characterized as having 57 58 repetitive episodes of giving way, decreased self-reported function, ongoing symptoms such as pain or weakness, and recurrent ankle sprains for at least one year following the initial LAS.⁶ 59 60 Individuals with CAI will all have primary tissue injury to the lateral ankle ligament(s), however, impairments will be unique to each individual.³ Hertel and Corbett² categorize these impairments 61 found within individuals with CAI as motor-behavioral, sensory-perceptual, and 62 pathomechanical impairments. Motor-behavioral impairments that often present as aberrant 63 biomechanical patterns during functional and dynamic movements have been well-documented 64 in CAI.³ 65 Several altered gait characteristics have been observed during walking in individuals with 66

CAI compared to individuals with a history of no LAS and individuals that have a history of 67 LAS but who return to pre-injury health status (termed copers). Individuals with CAI often 68 display greater ankle inversion throughout the gait cycle^{7–9} which may coincide with a lateral 69 deviation in the center of pressure (COP)^{10,11} and increased plantar pressure along the lateral 70 column of the foot during walking.^{10,12} This biomechanical profile of gait is associated with an 71 elevated risk of LAS and may contribute to the earlier onset of ankle posttraumatic osteoarthritis 72 (PTOA) in individuals with CAI.^{13–15} When the location of COP approaches the lateral boundary 73 74 of the foot, it places the ankle in a position similar to that of a LAS and may lead to recurrent

sprains. Similarly, increased subtalar joint supination at touch down during a side shuffling task
simulation has been shown to increase the occurrence of LAS while decreased supination
decreased the occurrence of LAS.¹⁶ Unfortunately, this position can also result in abnormal stress
distribution throughout the talar cartilage, thereby influencing the development of ankle
PTOA.^{17,18} Therefore, it is crucial to restore gait patterns in individuals with CAI to maintain
long-term joint health of the ankle.

Various approaches have been utilized to address these abnormal gait patterns in 81 individuals with CAI, including traditional rehabilitation techniques such as strength and balance 82 training,^{19,20} as well as targeted gait training strategies involving the use of devices or 83 biofeedback methods.^{21,22} While traditional rehabilitation strategies are successful at improving 84 strength and balance when trained, they have not shown evidence of improving gait 85 biomechanics.^{19,20} Recently published critically appraised topics have evaluated the effectiveness 86 of taping and bracing,²³ neuromuscular training,²⁴ and gait biofeedback training²¹ for improving 87 gait impairments in individuals with CAI. Of the aforementioned interventions, only biofeedback 88 training showed efficacy at improving the specific gait pattern (i.e. lateralized COP) associated 89 with CAI.²¹ 90

Biofeedback training involves providing a stimulus (visual, auditory, haptic) to correct
unwanted movement patterns and appears to be effective at improving respective gait
biomechanical outcome measures (kinematics and plantar pressure).²¹ Another technique to
address gait alterations has been the implementation of gait training devices such as
destabilization devices^{25–27} and a custom gait training device using resistance bands.²⁸
Destabilization devices are worn by individuals to create an unstable surface under the foot with
the goal of improving neuromuscular control in patients with CAI that exhibit symptoms

associated with sensory-perceptual impairments such as perceived instability. Sensory-perceptual
impairments have been defined as how the individual senses or feels about the body, injury, or

100 themselves.³

Several gait training strategies have been investigated for improving aberrant 101 102 biomechanics in individuals with CAI, however, a systematic review of the literature with meta-103 analysis has yet to be conducted to synthesize this information and provide a synopsis on the 104 effectiveness of these gait training interventions in individuals with CAI. The purpose of this study was to systematically review the literature on the efficacy of gait training interventions 105 106 (devices, biofeedback) for improving altered gait biomechanics in individuals with CAI. 107 Methods 108 Search Strategy 109 This systematic review with meta-analysis was registered with the International Prospective Register of Systematic Reviews (PROSPERO, CRD4202XXXXXX) on 110 September 12, 2022. Preferred Reporting Items for Systematic Reviews and Meta-Analysis 111 112 (PRISMA) guidelines were followed while conducting this systematic review and metaanalysis.²⁹ A health sciences librarian was consulted for the development of a systematic search 113 of electronic databases. The search was performed in the online search engines PubMed, 114 115 CINAHL, SPORTDiscus, and MEDLINE from database inception through September 15, 2022, 116 using the following search terms ((Chronic ankle instability OR CAI OR functional ankle 117 instability OR recurrent ankle sprain) AND (gait training OR gait devices OR biofeedback OR 118 feedback) AND (biomechanics OR kinetics OR kinematics OR electromyography)). Searches

119 were filtered for English language and full-text available. Following the initial search, screening

120 of the literature and data extraction were completed. Two authors (XX, XX) independently

screened all titles, abstracts, and full-text records for eligible studies (Figure 1). If conflicts

122 existed, the authors discussed the study to reach consensus. If consensus was not achieved, a

third author (XX) was consulted. Manual reference list screening was performed to identify any

- additional studies.
- 125 Study selection criteria and quality assessment

126 Studies were included if they met the following criteria: 1) included individuals with CAI (as determined using the International Ankle Consortium guidelines),⁶ 2) a gait training 127 intervention was administered using devices or biofeedback methods, 3) outcome measures 128 129 included gait kinetics, kinematics, and/or muscle activity during walking, 4) study was published in peer-reviewed journal, and 5) full-text was published in English. Randomized controlled trails, 130 cross-over design, quasi-experimental design, and descriptive laboratory or field studies were 131 included. Studies were excluded if individuals with CAI were not included, interventions did not 132 involve gait training, biomechanical outcomes were not measured, not available in English, 133 and/or the full text was unavailable. 134

135 The Downs and Black quality assessment checklist was used to evaluate the included studies (Table 1).³⁰ The checklist consists of 27 questions within 5 sections (reporting, external 136 validity, internal validity, internal validity confounding [selection bias], and power) and was 137 designed to assess the methodological quality of randomized and nonrandomized comparative 138 studies.³⁰ Questions were scored as yes (1), no (0), or not applicable (/) with the exception of 139 140 question 5, which was scored as yes (2), partially (1), no (0), or not applicable with a maximum total score of 28 points.³⁰ Higher scores indicated higher methodological quality.³⁰ Two authors 141 142 (XX, XX) independently scored all included studies. If scores did not align, a third author (XX) 143 was consulted.

144 Data extraction

Study design, participant demographics, sample sizes, intervention type (device,
biofeedback), intervention length, and biomechanical outcome measures (kinematics, kinetics,
electromyography (EMG)) were extracted by one author (XX) for all included studies (Tables 24). Authors were contacted if values were not reported in the text or were presented as graphs.
When at least 3 studies reported on the same outcomes, the mean and standard deviations were
extracted for potential meta-analyses.

151 Data analysis

When 3 or more studies reported on the same outcome measure using consistent units or 152 units that could be derived for equivocal comparisons, meta-analyses were conducted. Meta-153 analyses were performed using a random-effects model in JASP software (JASP Team 2023, 154 version 0.17.2.1, University of Amsterdam, Netherlands) to compare differences before and 155 during administration of gait training for studies involving a single session for the following 156 variables: kinetics (COP gait line, peak pressure, contact area, contact time, pressure time 157 integral) and EMG (root mean square [RMS] amplitude pre-initial contact [IC] and post-IC). 158 Meta-analyses were considered statistically significant when p<.05. There were not enough 159 multi-session gait training studies for meta-analyses to be conducted on any variables. Meta-160 161 analysis ES and associated 95% confidence intervals were displayed using forest plots (Figures 162 2-4). Effect sizes (ES) and standard error of ES using the pooled standard deviation were 163 calculated to determine that magnitude of difference between time points (pre- vs. post-gait training) or between groups (gait training vs. no gait training). ES were interpreted as very small 164 (<0.20), small (0.21-0.39), medium (0.40-0.79), and large (>0.80).³¹ Heterogeneity was analyzed 165 using the I^2 test statistic and summarize the variation across studies due to difference rather than 166

167	chance as recommended by the Cochran Handbook for Systematic Reviews of Interventions. ³²
168	Interpretation of the I^2 test statistic used the following guidelines: 0-40% may not be important,
169	30-60% may represent moderate heterogeneity, 50-90% may represent substantial heterogeneity,
170	75-100% considerable heterogeneity. ³² When heterogeneity was considerable (I^2 >75%), studies
171	showing the same direction of effect were still considered appropriate for meta-analysis. ³³
172	Publication bias was assessed using funnel plots and associated Egger's regression test for
173	variables identified as statistically significant by the meta-analyses. Significant publication bias
174	was considered present when p<.05 for Egger's regression test. ³⁴
175	Results
176	Study Selection & Characteristics
177	Our initial search yielded 358 studies (Figure 1). Following duplicate removal, abstract
178	screening, and full-text review, 13 studies were included. ^{25–28,35–43} Of the studies included, 11
179	reported on kinetic outcome measures, ^{25,26,28,35,40,42,43} 3 reported on kinematic outcome
180	measures, ^{25,39,41} and 7 reported on muscle activity outcome measures. ^{25–28,35,36,39} Of the studies
181	included, 5 utilized a gait training device such as a destabilization sandal or boot ^{25–28,36} and 8
182	utilized a form of biofeedback (visual, auditory, haptic). ^{35,37–43} Summaries of the study
183	characteristics, outcome measures, and results for kinetics, kinematics, and muscle activity are
184	presented in tables 2-4 respectively.
185	Methodological Quality Assessment
186	Downs and Black scores for the included studies ranged from 16-25 points out of a
187	maximum 28 points. The 3 studies with randomized controlled trial study designs had the highest
188	overall scores with a range of 24 points ⁴³ to 25 points. ^{25,39} Reviewers scored all studies "yes" or

189 "not applicable" to all questions within the reporting section of the checklist with the exception

190	of if adverse events that may impact the intervention were reported (Question [Q] 8). For the
191	external validity section, all studies scored "yes" for subjects representative of the population
192	they were recruited from (Q11) and subjects who were prepared to participate represented the
193	population from which they were recruited (Q12). All studies scored "no" for if staff, places, and
194	facilities where patients were treated were representative of the treatment majority of the patients
195	received (Q13). The studies were scored "no" for Q13 because the gait training methods
196	employed in the research studies were not representative of treatments in common use in clinical
197	practice settings for individuals with CAI. Additionally, gait training visits were conducted under
198	the supervision of a research team using unique equipment for administering gait training that is
199	not currently available to clinicians or individuals with CAI. When considering internal validity
200	subscale, no studies made an attempt to blind the study subjects to the intervention (Q14). In the
201	randomized controlled trials only, ^{25,39,43} attempts were made to blind the individual measuring
202	the main outcome measures of the intervention (Q15). All studies scored "yes" or "not
203	applicable" for the remaining internal validity questions (Q16-20). When considering the internal
204	validity – confounding (selection bias) subscale, all studies subjects in intervention groups were
205	recruited from the same population (Q21) and all studies accounted for subjects lost to follow-up
206	(Q26). All randomized controlled trials ^{25,39,43} randomized subjects into intervention groups
207	(Q23), randomization was concealed (Q24), and adequate adjustments for confounding in the
208	analyses for main findings were made (Q25). Only 3 studies ^{26,27,39} reported a sample size
209	estimate needed to meet the power calculation requirement for Q27.

210 *Heterogeneity*

Heterogeneity ranged from 0-40% and was interpreted to be not important for the 10%

212 increments of the COP gait line at all time points (range 0 to 35.9%), contact time for the medial

216	Publication Bias Assessment
217	Funnel plots and associated Egger's regression test results for the meta-analyses are
218	reported in the supplemental figure. Publication bias was present for the location of COP during
219	0-10% of the stance phase ($p=.015$), for peak pressure in the lateral midfoot ($p<.001$), and for the
220	PTI in the lateral midfoot (p<.001). There were no other significant findings for publication bias
221	for any other measures included in our meta-analyses.
222	Gait Training Approaches
223	Five studies utilized gait training devices ^{25–28,36} and 8 studies utilized biofeedback ^{35,37–43}
224	for gait training. Among the gait training device studies, 2 used destabilization boots and
225	sandals, ^{25,27} 1 used a wearable multi-axis destabilization device, ²⁶ and 2 used a custom-built gait
226	training device with resistance bands. ^{28,36} Among the biofeedback gait training studies, 3 used
227	visual biofeedback, ^{37,39,42} 2 used auditory biofeedback, ^{35,43} and 3 used haptic biofeedback. ^{38,40,41}
228	For the visual biofeedback, 1 study used a shoe mounted cross-line laser with instructions to
229	"walk in a manner in which the vertical laser line aligns with the piece of tape on the wall," ⁴² 1
230	study used real-time 2D video from the posterior aspect of the treadmill with instructions to
231	"walk in a manner where you can no longer view the outside or inside of your foot on the
232	television screen while you walk,"37 and 1 study used a custom real-time display of ankle
233	inversion angles that turned red for steps with ankle inversion above the set threshold (too much
234	inversion) or green for steps within the desired range for ankle inversion with instructions to
235	"avoid walking on the outside of your foot so as not to exceed the inversion threshold." ³⁹ For the

forefoot (37.0%), peak pressure for the hallux (17.6%), and pressure time integral for the lateral

heel (3.8%) and hallux (3.6%). Heterogeneity was >75% and was interpreted as considerable for

peak pressure and pressure time integral in the lateral midfoot (87.4% and 90.8% respectively).

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auditory biofeedback, 2 studies used a custom device that was created to set a pressure threshold
under the lateral aspect of the foot and provide an auditory tone when the participant's vertical
force exceed the set threshold.^{35,43} For the haptic biofeedback, 3 studies used a custom device
similar to the auditory biofeedback studies, however, instead of delivering an auditory tone,
vibration was provided on the lateral malleolus of the test limb when the participant's vertical
force exceeded the set threshold under the lateral aspect of the foot.^{38,40,41}

242 Kinetic Outcomes

Eleven studies examining kinetic outcomes met the inclusion criteria for this systematic 243 review (table 2). Six studies reported on the COP gait line.^{26,28,36,40,42,43} The COP gait line was 244 defined as the location of COP from most medial border of the foot at 10% increments in 5 245 studies^{26,28,36,42,43} and the location of COP in the lateral-medial direction from the position of the 246 247 marker at the 5th metatarsal head with the foot modeled as a rectangle at 10% increments in 1 study.⁴⁰ Of these studies, 4 were single session^{26,28,40,42} and results were pooled for meta-analyses 248 (Figure 2). The meta-analyses revealed there were small to large medial shifts in the location of 249 COP at each 10% increment from 0-90% (ES range: -0.35 to -0.82, I^2 range: 0 to 35.911, p-value 250 range: <.001 to .041, Egger's regression p-value range: .015 to .125) for the COP gait line. Seven 251 studies reported on traditional plantar pressure measures (contact area, contact time, peak 252 253 pressure, pressure time integral [PTI], time to peak pressure) and results were pooled for metaanalyses (Figure 3).^{26,28,35–37,42,43} Contact area was defined as how large of an area of each region 254 255 of the foot was in contact with the ground during the stance phase and was measured in centimeters squared (cm²).^{28,35–37,42} Contact time was defined as how much time each region of 256 257 the foot was in contact with the ground during the stance phase and was measured in milliseconds (ms).^{28,35–37} Peak pressure was defined as the highest amount of pressure in a given 258

259	region of the foot during the stance phase of gait and was measured in kilopascals (kPa). ^{26,28,35–}
260	^{37,42,43} PTI was defined as the total plantar pressure applied to a specific region of the foot
261	multiplied by the time spent in the stance phase of gait and was measured in kilopascals
262	multiplied by seconds (kPa*s). ^{28,35–37,42} Time to peak pressure was defined as the % of stance
263	when peak pressure occurred for the specified region of the foot. ^{28,35,36} Meta-analyses revealed
264	that contact time was significantly decreased in the medial forefoot (ES: -0.43 [-0.86,0.00],
265	I^2 =36.997, p=.049, Egger's regression p=.260). Peak pressure was significantly increased in the
266	hallux (ES: 0.59 [0.21,0.96], I^2 = 17.624, p=.002, Egger's regression p=.156) and significantly
267	decreased in the lateral midfoot (ES: -1.18 [-2.24,-0.12], I^2 = 87.438, p=.029, Egger's regression
268	p<.001). Pressure time integral was increased in the hallux (ES. 0.63 [0.30,0.97], I^2 = 3.556,
269	p<.001, Egger's regression p=.144) and decreased in the lateral heel (ES: -0.33 [-0.66,0.00], I^2 =
270	3.775, p=.050, Egger's regression p=.066) and lateral midfoot (ES: -1.22 [-2.43,0.00], I^2 =
271	90.757, p=.049, Egger's regression p<.001). There were no other significant differences from the
272	meta-analyses for any other kinetic parameters. Two studies reported on internal joint moments
273	and found no significant differences after gait training. ^{25,39} Only one study reported on impact
274	peak, time to impact peak, impact loading rate, propulsive peak, time to propulsive peak,
275	propulsive loading rate, ankle joint contact force peak, ankle joint contact force impulse, and
276	ankle joint contact force loading rate. ³⁸

277 Kinematic Outcomes

Three studies examining kinematic outcome measures met the inclusion criteria for the systematic review (Table 3).^{25,39,41} Three studies measured 3-Dimensional (3D) ankle joint angles (°)^{25,39,41} and 2 of those studies measured 3D joint angles at the knee and hip.^{25,39} All studies reported 3D ankle kinematics at IC and throughout the loading phase (first 10% of

282 stance), however, only one study was a single session gait training study so meta-analyses were not performed.⁴¹ Decreased ankle inversion during the loading response was reported by 2 283 studies^{39,41} and 1 study found no differences in ankle inversion.²⁵ Only one study reported on 284 285 hindfoot and forefoot joint angles and found increased forefoot abduction during the loading 286 phase in the laboratory and real world settings and increased forefoot abduction during the loading phase in the laboratory setting.⁴¹ Two studies reported on ankle, knee, and hip 287 kinematics throughout the stride cycle (0-100%).^{25,39} One study reported increased external 288 rotation at the knee during terminal swing³⁹ with a medium ES while the other study found no 289 significant differences.²⁵ Significant differences were not identified by either study for hip joint 290 angles.^{25,39} 291

292 Muscle Activity Outcomes

293 Seven studies measured muscle activity using and met the inclusion criteria for the systematic review (Table 4).^{25–28,35,36,39} Of the included studies, 4 reported EMG RMS 294 amplitudes for the 50-200ms pre-IC and 200ms post-IC, 2 studies reported EMG RMS 295 amplitudes throughout the stride cycle (0-100%),^{25,39} and 1 study reported EMG RMS 296 amplitudes during the stance phase (0-100%).³⁶ Meta-analyses were conducted for the EMG 297 RMS amplitudes pre-IC and post-IC for the tibialis anterior, fibularis longus, and gluteus medius 298 299 muscles. During the 200ms post-IC, muscle activity was increased during gait training for the fibularis longus muscle (ES: 0.83 [0.43, 1.22], $I^2=0$, p<.001, Egger's regression p=.986) (Figure 300 301 4). There were no other significant differences identified by the meta-analyses for any other muscle activity parameters. Prior to IC, 2 studies reported increased fibularis longus activity^{27,28} 302 with large ES and 2 reported no differences for fibularis longus activity.^{26,35} During the stance 303 phase, 1 study reported increased fibularis longus activity with medium to large ES,³⁶ while 304

another study reported decreased fibularis longus activity with large ES,²⁵ and a third study

306 reported no significant differences.³⁹ For the tibialis anterior muscle activity, 1 study reported

increased activity pre-IC 26 with a large ES, while 3 studies reported no significant

308 differences.^{27,28,35} One study reported decreased gluteus medius activity during late stance³⁶ with

309 medium to large ES and 2 studies reported no significant differences.^{25,39}

310 Discussion

311 This systematic review with meta-analysis identified 13 studies that measured biomechanical outcomes before and after gait training in individuals with CAI. We categorized 312 biomechanical outcome measures into kinetics, kinematics, and muscle activity. Among the 313 studies included, 11 measured kinetics, 3 measured kinematics, and 7 measured muscle activity 314 making meta-analyses possible for several outcome measures. Gait training techniques included 315 wearing destabilization devices,^{25–27} using a custom gait training device with resistance 316 bands,^{28,36} or using biofeedback including auditory,^{35,43} visual,^{37,39,42} or haptic^{38,40,41} biofeedback 317 modes aimed to improve various biomechanical outcome measures. Based on the results from 318 319 the meta-analyses, a single session of gait training improved COP location, reduced lateral plantar pressure, and increased muscle activity in the fibularis longus muscle during the 200ms 320 post-IC. Targeted gait training improved corresponding gait biomechanics in almost all studies. 321 Few studies required multiple sessions of gait training^{25,36,39,43} and longer term effects of gait 322 training were not well documented with the longest follow-up time being 1-week.⁴³ 323

324 *Methodological Quality*

Studies included in our systematic review and meta-analysis were critically appraised
using the Downs & Black scoring system. Study quality using the Downs & Black scoring has
previously been categorized as excellent (26-28), good (20-25), fair (15-19), and poor (<15).⁴⁴

328 The scores of the included studies ranged from 16-25 points out of a possible 28 points 329 demonstrating fair to good methodological quality (Table 1). The randomized controlled trials 330 had the highest methodological quality (24-25 points) followed by the quasi-experimental trial 331 (17 points) and descriptive laboratory study designs (16-18 points). Studies did not satisfy all 332 criteria because information was not included or explicitly stated within the published 333 manuscript and therefore could not earn points for that question. For the reporting section, the 334 majority of studies scored "yes" or "not applicable" for all questions, except for if adverse events were reported. None of the included studies reported or mentioned any adverse events associated 335 336 with gait training which may suggest that the gait training techniques employed by the studies are not high risk for the given population. The included studies scored "yes" to all questions in 337 the external validity section except for if the staff, places, and facilities were representative of 338 339 treatments patients receive. This is not surprising as all studies took place in a laboratory setting and utilized techniques that are not currently available to most practicing clinicians. 340 Additionally, gait training methods explored in the research studies were not representative of 341 342 current treatments used in treating individuals with CAI. While study methods do not reflect current treatment methods in the clinical setting, these studies provide the foundational evidence 343 to support that gait biomechanics in individuals with CAI may be improved through various gait 344 345 training methods. To improve external validity, future studies should explore gait training 346 methods that can be easily implemented in clinical practice for individuals with CAI. Scores 347 were high within the internal validity section, however, no studies blinded subjects to the 348 intervention. This would be a considerable challenge given the primary modes of gait training 349 involve wearing devices or responding to some form of immediate biofeedback. When 350 considering the confounding or selection bias within the next internal validity section, scores

were low. Most studies did not report the timeframe in which participants were recruited, did not
randomize participants into intervention groups and did not conceal randomization apart from the
randomized controlled trials.

354 *Heterogeneity*

355 For peak pressure and PTI in the lateral midfoot, heterogeneity was considerable (87.4% 356 and 90.8% respectively), however, all studies showed the same direction of effect and were 357 therefore still considered appropriate for inclusion in the meta-analyses. Higher levels of heterogeneity may indicate that studies are measuring different underlying effects or there are 358 359 methodological differences between the studies. Upon further inspection of the individual studies included in the meta-analyses for peak pressure and PTI.^{28,35,37,42} all studies utilized the Pedar-X 360 plantar pressure system to measure and analyze plantar pressure outcomes, however, the gait 361 training interventions varied greatly between studies. For example, Donovan et al.³⁵ utilized an 362 auditory biofeedback device placed under the 5^{th} metatarsal, Feger and Hertel²⁸ created a custom 363 gait training device using resistance bands, Ifarraguerri et al.³⁷ projected a live video of a 364 posterior view of the foot in front of the treadmill, and Torp et al.⁴² placed a crossline laser on 365 top of the foot. The studies utilizing auditory feedback and the custom gait training device found 366 significant reductions in peak pressure and PTI in the lateral midfoot while the studies using the 367 live video and crossline laser found no significant differences while receiving gait training. It is 368 369 possible that the substantial variations in gait training methods contributed to the considerable 370 heterogeneity found by the meta-analyses.

371 Publication Bias Assessment

Publication bias was evaluated using funnel plots and Egger's regression tests in our
meta-analyses. Notably, significant publication bias was detected regarding the location of the

374 COP from 0-10% of the stance phase, peak pressure in the lateral midfoot, and PTI in the lateral

375 midfoot. These findings indicate a potential overstatement of results pertaining to these measures

376 within our meta-analyses. Such bias may skew the meta-analysis ES upward, thus potentially

377 inflating the results and inaccurately suggesting a stronger ES that may be attributable to random

378 chance. The detection of publication bias suggests there may be an overrepresentation of studies

379 reporting positive outcomes for these measures. This bias may distort the overall findings,

leading to an inflated perception of the ES and potentially resulting in misleading conclusions.

381 *Kinetics*

Kinetic variables were the most frequently reported by studies included in this systematic 382 review. The COP gait line is described as the mediolateral location of COP at 10% increments 383 during the stance phase¹⁰ and was the most frequently reported kinetic variable. All 384 studies^{26,28,36,42,43} utilized the Pedar-X plantar pressure system to measure and analyze the 385 location of COP except for Migel and Wikstrom⁴⁰ Gait training strategies to target the COP gait 386 line included a custom gait training device with resistance bands,^{28,36} multi-axis destabilization 387 devices,²⁶ visual biofeedback,⁴² haptic biofeedback,⁴⁰ and auditory biofeedback.⁴³ The meta-388 analyses revealed that from 0-90% of the stance phase, gait training shifted the COP gait line 389 medially while participants received gait training. The pooled ES ranged from -0.35 to -0.82 390 391 throughout the stance phase suggesting small to large improvements. Studies involving multiple gait training sessions^{36,43} tended to show greater medial shifts in the COP gait line as seen with 392 393 the larger MD following gait training sessions (table 2). The medium to large medial shift in COP is considered beneficial because when the center of gravity approaches or exceeds the 394 lateral boundary of the foot, an episode of giving way or LAS may occur.¹⁴ Various gait training 395

strategies were effective at reducing laterally deviated COP and should be implemented whenindicated for individuals with CAI.

398 Traditional plantar pressure measures (contact area, contact time, peak pressure, PTI, 399 time to peak pressure) were often reported for nine specified regions of the foot including: 400 medial heel, lateral heel, medial midfoot, lateral midfoot, medial forefoot, central forefoot, lateral forefoot, hallux, and toes 2-5 in 7 studies.^{26,28,35–37,42,43} Gait training strategies to target the 401 traditional plantar pressure measures included a custom gait training device with resistance 402 bands,^{28,36} multi-axis destabilization devices,²⁶ visual biofeedback,^{37,42} and auditory 403 biofeedback.^{35,43} Generally speaking, traditional plantar pressure measures were reduced in the 404 lateral aspect of the foot and pressure shifted medially which is the desired outcome for gait 405 training in individuals with CAI. Individual studies reported decreased contact area for the lateral 406 midfoot^{28,35} or increased contact area in the medial midfoot^{36,42} suggesting a medial shift in 407 pressure area may exist following gait training 408 Peak pressure was considered the maximum loading in an area under the foot.^{26,28,35–} 409

^{37,42,43} The meta-analyses revealed decreased pressure in the lateral midfoot with a large ES and a 410 medium increase in peak pressure for the hallux. Increased peak pressure for total foot was 411 reported by 2 studies.^{28,35} Although not investigated among patients with CAI, the overall 412 413 increase in peak pressure may be a beneficial adaptation when regarding PTOA. Studies have 414 found that greater mechanical loading during walking is associated with less type II collagen turnover among patients who underwent anterior cruciate ligament reconstruction.^{45,46} Similar to 415 peak pressure, PTI was described as the total amount of pressure for a specific region of the foot 416 multiplied by the time spent in stance.^{28,35–37,42,43} The meta-analyses revealed that PTI decreased 417 418 in the lateral heel and lateral midfoot and increased in the hallux again suggesting a shift from

lateral to medial plantar pressure. The results for the peak pressure and PTI in the lateral midfoot
should be interpreted with caution. Considerable heterogeneity and significant publication bias
were identified for the meta-analyses for these outcomes and suggest that the larger ES for these
outcomes may be due to chance rather than an actual observed change. Future studies involving
larger sample sizes assessing the effects of gait training on these plantar pressure outcome
measures are therefore warranted.

425 The results from our meta-analyses suggest that several plantar pressure measures are significantly improved by various gait training methods involving devices or biofeedback 426 techniques. Many individuals with CAI demonstrate increased plantar pressure along the lateral 427 column of the foot which may be associated with an elevated risk of LAS and could contribute to 428 the earlier onset of ankle PTOA in individuals with CAL^{13–15} Shifting the pressure medially 429 reduces the risk of the COP approaching the lateral boundary of the foot potentially resulting in 430 an LAS. This altered ankle position can also result in abnormal stress distribution throughout the 431 talar cartilage, thereby influencing the development of ankle PTOA.^{17,18} Therefore, it is crucial to 432 restore gait patterns in individuals with CAI to maintain long-term joint health of the ankle 433 which appears to be possible through the utilization of gait training. 434

435 *Kinematics*

Kinematics were the least reported outcome measures with only 3 studies meeting the
inclusion criteria for the systematic review.^{26,28,35–37,42,43} Gait training strategies to target the
kinematic measures included destabilization devices,²⁵ visual biofeedback³⁹ and haptic
biofeedback.⁴¹ Two studies found that gait training with biofeedback (haptic and visual) reduced
ankle inversion by 2.5-7.3^{o39,41} while 1 study using destabilization devices found no significant
changes in ankle inversion following gait training but found increased ankle dorsiflexion by 5.4°

during mid-late stance.²⁵ Of the studies included, only 1 specifically targeted the reduction of 442 ankle inversion as part of the gait training protocol.³⁹ Because only 3 studies utilizing gait 443 training to improve biomechanics in individuals with CAI measured kinematic outcomes, it is 444 445 difficult to understand the utility of gait training for improving ankle kinematics at this time, 446 however, it is likely that the medial shift in the COP gait line and additional plantar pressure 447 outcome measures could be associated with shifting from an inverted to everted ankle position. 448 Walking with the foot in an everted position has been shown to create more contact under the medial aspect of the foot and thus the COP was located on the medial aspect of the foot.⁴⁷ Future 449 gait training studies for individuals with CAI measuring kinematic outcomes should consider 450 techniques targeting ankle inversion specifically. 451

452 Muscle Activity

Muscle activity was measured in 7 studies^{25–28,35,36,39} using EMG and RMS amplitude 453 was reported for all included studies, however, the timing during the stride cycle that data were 454 reported for differed among studies making meta-analyses possible only for short time periods 455 pre-IC and post-IC. ^{26,28,35–37,42,43} Gait training strategies were not specifically used to target 456 muscle activity, however, several studies measured muscle activity as a primary outcome 457 measure and included a custom gait training device with resistance bands,^{28,36} destabilization 458 devices,^{25–27} visual biofeedback,³⁹ and auditory biofeedback.³⁵ Our meta-analyses revealed a 459 460 large increase in fibularis longus activity during the 200ms post-IC while receiving gait training. 461 Increased fibularis longus activity immediately following IC during the loading response may be beneficial in contributing to increased ankle stability and the medial shift in plantar pressure.^{10,48} 462 463 Individuals without a history of LAS have been shown to activate their fibularis longus during midstance to assist with pronation and stabilizing the first ray during propulsion.⁴⁹ 464

465	There were several limitations that should be considered when interpreting the results of
466	this study. Individual study sample sizes were relatively small and only included 10-27
467	participants. Results from these studies should be interpreted with caution and further research is
468	needed in this area. The timing in which biomechanical outcomes were measured varied among
469	studies. Several studies measured gait outcomes while participants were wearing devices ^{26,28} or
470	receiving biofeedback ^{37,38,42,43} while other studies measured outcomes after gait training had
471	commenced. ^{25–27,36,39–41,43} Gait training protocols differed substantially between studies. For
472	example, studies in the visual biofeedback category involved a variety of techniques including
473	projecting real time ankle kinematics displayed in front of the treadmill, ³⁹ real time video of the
474	posterior aspect of the ankle, ³⁷ and using a cross-line laser attached to the dorsal aspect of the
475	foot. ⁴² In addition, the number of gait training sessions implemented for each study protocol
476	ranged from a single session up to 12 total sessions which may influence the effects of gait
477	training on biomechanical outcomes. Lastly, the gait training methodology utilized by many
478	studies is not currently clinically accessible which makes implementation unrealistic for athletic
479	trainers or other health care professionals treating individuals with CAI. Future studies should
480	consider gait training techniques that would be feasible for clinical implementation.
481	There are several future directions to consider for gait training implementation for
482	individuals with CAI. New gait training strategies should attempt to transition concepts from
483	laboratory-based interventions to strategies using minimal or no equipment to increase the
484	feasibility of implementation in the clinical setting. Future studies should also consider assessing
485	long-term outcomes, dosage, measures of joint health, and the risk reduction of subsequent LAS
486	associated with gait training. While this study has established that gait training can be utilized to
487	improve a variety lower extremity gait biomechanics immediately and for a short duration (up to

488 1-week), long-term outcomes are not yet understood. Another component of gait training to 489 consider are the total number of gait training sessions and the length of sessions needed to 490 improve and maintain desired gait changes. This information may be useful in determining if 491 additional sessions are needed as a booster or refresher following the cessation of gait training 492 programs to maintain desired gait changes. The overarching goal of gait training should not only 493 be to improve biomechanics, but also to improve ankle joint health and reduce the risk of future 494 LAS. Future research should address these critical areas to continue facilitating gait training and its broader adoption in clinical practice for patients with CAI. 495 496 Conclusion

Gait training protocols included in the systematic review utilized devices or biofeedback 497 to effectively improve lower extremity biomechanics in individuals with CAI. These 498 499 interventions resulted in notable improvements such as medial shifts in plantar pressure, decreased ankle inversion, and increased fibularis longus activity which may be associated with 500 reducing the risk of LAS and development of ankle PTOA.^{17,18} It is worth noting that current gait 501 502 training strategies may present practical challenges within the clinical setting. Therefore, future research endeavors should investigate alternative techniques that are more accessible for clinical 503 implementation. It is critical to restore gait patterns in individuals with CAI which appears to be 504 possible through the utilization of gait training. 505

506 **Conflicts of interest**

507

The authors have no conflict of interest to declare in relation to this article.

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Authors	Participants	Study Information	Main Findings
Donovan et al.	26 CAI (13 control,	Design: Randomized controlled trial	<i>vGRF (N/kg):</i> No significant differences
	13 intervention)	Gait Training: 12 sessions with device	Internal joint moments (Nm/kg): No significant
		(destabilization boot and sandal)	differences
		<i>Outcome Measures:</i> vGRF, internal joint moments	
		<i>Data Collection Timepoints:</i> Baseline, 2-7 days after last gait training session	
Donovan et al.	10 CAI	Design: Descriptive laboratory study	<i>Contact area</i> (<i>cm</i> ²): Decreased in lateral midfoot (MD=-
			4.3, ES=-1.3) and toes 2-5 (MD=-2.1, ES=-0.7)
		Gait Training: I session with biofeedback (auditory)	Contact time (ms): No significant differences
		Outcome Measures: Contact area contact	Peak pressure (kPa). Decreased in lateral midfoot
		time, peak pressure, PTI, time to peak pressure	(MD=-52.8, ES=-2.8), central forefoot (MD=-29.8, ES=- 1.4), and lateral forefoot (MD=-57.8, ES=-2.4), increased at hallux (MD=91.7, ES=1.0)
		Data Collection Timepoints: Baseline, while receiving biofeedback	<i>PTI (kPa*s):</i> Decreased at lateral midfoot (MD=-28.4, ES=-3.1) and lateral forefoot (MD=-29.1, ES=-2.6), increased for hallux (MD=31.3, ES=1.1) and total foot (MD=18.6, ES=1.0)
			<i>Time to peak pressure (% of stance):</i> Reached earlier in lateral midfoot (MD=-15.9, ES=-1.0)

Table 1. Summary of articles related to kinetic outcome measures. All results are reported in comparison to baseline values.

Feger & Hertel 10 CAI

Design: Descriptive laboratory study

Gait Training: 1 session with device (novel gait trainer with resistance bands)

Outcome Measures: COP gait line, contact area, contact time, peak pressure, PTI, time to peak pressure

Data Collection Timepoints: Baseline, while using device

Design: Quasi-experimental trial

Gait Training: 5 sessions with device (novel gait trainer with resistance bands)

Outcome Measures: COP gait line, contact area, contact time, peak pressure, PTI, time to peak pressure *COP gait line (mm):* Medial shift from 0-100% of stance phase: 0-10% (MD=-4.7, ES=-1.7), 11-20% (MD=-3.8, ES=-1.1), 21-30% (MD=-3.6, ES=-0.9), 31-40% (MD=-4.6, ES=-1.0), 41-50% (MD=-5.4, ES=-1.2), 51-60% (MD=-6.4, ES=-1.4), 61-70% (MD=-7.0, ES=-1.6), 71-80% (MD=-7.1, ES=-1.6), 81-90% (MD=-6.4, ES=-1.5), 91-100% (MD=-5.2, ES=-1.2)

Contact area (cm²): Decrease in lateral midfoot (MD=-0.8, ES=-0.5)

Contact time (ms): No significant differences

Peak pressure (kPa): Decreased in lateral midfoot (MD=-29.8, ES=-1.5) and lateral forefoot (MD=-27.4, ES=-0.9), increased at lateral heel (MD=18.2, ES=1.0), medial heel (MD=23.2, ES=1.4), hallux (MD=72.9, ES=0.9), and total foot (MD=52.2, ES=0.7)

*PTI (kPa*s):* Decreased in lateral midfoot (MD=-13.8, ES=-1.4) and lateral forefoot (MD=-9.8, ES=-0.7) increased in medial forefoot (MD=7.4, ES=0.5), hallux (MD=22.3, ES=1.0), and total foot (MD=19.3, ES=0.9)

Time to peak pressure (% of stance): Occurred earlier in lateral midfoot (MD=-13.1, ES=-0.7)

COP gait line (mm): Medial shift from 11-100% of stance phase: 11-20% (MD=-1.6, ES=-0.4), 21-30% (MD=-2.8, ES=-0.8), 31-40% (MD=-4.3, ES=-1.1), 41-50% (MD=-6.5, ES=-1.1), 51-60% (MD=-7.8, ES=-2.0), 61-70% (MD=-6.7, ES=-1.8), 71-80% (MD=-5.2, ES=-1.5), 81-90% (MD=-4.7, ES=-1.5), 91-100% (MD=-5.3, ES=-1.5)

Contact area (*cm*²): Increase in medial midfoot

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			(MD=3.0, ES=0.4)
		<i>Data Collection Timepoints:</i> Baseline, 24-72 hours after last gait	<i>Contact time (ms):</i> No significant differences
		training session	<i>Peak pressure (kPa):</i> Increased at hallux (MD=15.3, ES=0.4)
			<i>PTI (kPa*s):</i> Increased in medial forefoot (MD=4.4, ES=0.3)
			<i>Time to peak pressure (% of stance):</i> No significant differences
Ifarraguerri et al	26 CAI	Design: Descriptive laboratory study	Contact area (cm ²): No significant differences
		<i>Gait Training:</i> 1 session with biofeedback (visual)	<i>Contact time (ms):</i> No significant differences
		Outcome Measures:	<i>Peak pressure (kPa):</i> Decreased in medial forefoot (MD=-15.7, ES=-0.3)
		PTI	<i>PTI (kPa*s)</i> : Decreased in medial forefoot (MD=-2.3, ES=-0.1)
		<i>Data Collection Timepoints</i> : Baseline, while receiving biofeedback	
Jang et al.	10 CAI	Design: Descriptive laboratory study	Impact peak vGRF (N/BW): No significant differences
		<i>Gait Training:</i> 1 session with biofeedback (haptic)	<i>Time to impact peak vGRF (s):</i> No significant differences
		<i>Outcome Measures:</i> vGRF (impact peak, time to impact peak, impact leading rate, propulsive peak, time	<i>Impact loading rate vGRF (BW/s):</i> No significant differences
		to propulsive peak, propulsive loading rate), ankle JCF (peak, impulse, loading rate)	<i>Propulsive peak vGRF (N/BW):</i> Decreased during early (MD=-0.04, ES=-0.6) and late periods (MD=-0.04, ES=-0.5)

Data Collection Timepoints:

Baseline, while receiving biofeedback (early period = minute 1-2, late period = minute 9-10 of receiving biofeedback) *Time to propulsive peak vGRF (s):* Decreased during early (MD=-0.02, ES=-0.4) and late periods (MD=-0.02, ES=-0.5)

Propulsive loading rate vGRF (BW/s): Decreased during early period (MD=-0.24, ES=-0.4)

Ankle JCF peak (N/BW): Decreased during early period (MD=-0.24, ES=-0.4)

Ankle JCF impulse (BW*s): Decreased during early (MD=-0.09, ES=-0.6) and late periods (MD=-0.14, ES=-

Ankle JCF loading rate (BW/s): No significant differences

COP gait line (mm):

Wearing device: Medial shift from 11-60% of stance phase: 11-20% (MD=-5.7, ES=-1.1), 21-30% (MD=-6.3, ES=-1.2), 31-40% (MD=-6.2, ES=-1.2), 41-50% (MD=-5.8, ES=-1.1), 51-60% (MD=-5.0, ES=-0.9)

Post gait training: No significant differences

Peak pressure (kPa %):

Wearing device: Decrease in lateral midfoot (MD=-21.5, ES=-1.3), lateral forefoot (MD=-22.4, ES=-1.2), and central forefoot (MD=-17.5, ES=-1.0)

Post gait training: No significant differences

Knuckles et al. 12 CAI

Data Collection Timepoints: Baseline, while wearing device, immediately after devices removed

Outcome Measures: COP gait line, peak

Design: Descriptive laboratory stud

Gait Training: 1 session with dev

(multi-axis destabilization device)

pressure

Koldenhoven et al	27 CAI	Design: Randomized controlled trial	Internal joint moments (Nm/kg): No significant
ct di.	intervention)	<i>Gait Training:</i> 8 sessions with biofeedback (visual)	differences
		<i>Outcome Measures:</i> 3D internal joint moments for ankle, knee, hip throughout for 0-100% of stride cycle	
		<i>Data Collection Timepoints:</i> Baseline, 24-72 hours after last gait training session	
Migel et al.	19 CAI	Design: Descriptive laboratory study with repeated measures	COP gait line (mm):
		<i>Gait Training:</i> 2 sessions (1 laboratory, 1 real world) with biofeedback (haptic)	Medial shift from 0-90% of stance phase: 0-10% (MD=- 3.6, ES=-0.4), 11-20% (MD=-4.3, ES=-0.4), 21-30% (MD=-4.6, ES=-0.5), 31-40% (MD=-5.1 ES=-0.6), 41- 50% (MD=-5.1, ES=-0.7), 51-60% (MD=-4.2, ES=-0.7),
		<i>Outcome Measures</i> : COP gait line <i>Data Collection Timepoints</i> . Baseline, immediately after gait training, 5-minutes after gait training.	61-70% (MD=-2.8, ES=-0.6), 71-80% (MD=-1.7, ES=- 0.4), 81-90% (MD=-1.6, ES=-0.3)
			<i>Laboratory 5-minutes post:</i> Medial shift from 21-90% of stance: 21-30% (MD= -3.3, ES=-0.4), 31-40% (MD=-5.1, ES=-0.6), 41-50% (MD=-5.1, ES=-0.7), 51-60% (MD=-3.8, ES=-0.6), 61-70% (MD=-2.8, ES=-0.6), 71-80% (MD=-2.2, ES=-0.4), 81-

90% (MD=-1.6, ES=-0.3)

Real world immediately post:

Medial shift 0-70% of stance phase: 0-10% (MD=-6.0, ES=-0.8), 11-20% (MD=-7.3, ES=-0.8), 21-30% (MD=-

7.8, ES=-1.1), 31-40% (MD=-8.3, ES=-1.0), 41-50% (MD=-8.2, ES=-1.1), 51-60% (MD=-6.6, ES=-1.0), 61-70% (MD=-4.2, ES=-0.7), 71-80% (MD=-2.3, ES=-0.4)

Real world 5-minutes post:

Medial shift 0-60% of stance: 0-10% (MD=-4.7, ES=-0.5), 11-20% (MD=-5.6, ES=-0.6), 21-30% (MD=-6.1, ES=-0.8), 31-40% (MD=-6.5, ES=-0.7), 41-50% (MD=-5.9, ES=-0.7), 51-60% (MD=-4.1, ES=-0.6)

Torp et al. 26 CAI

Design: Descriptive laboratory study

Gait Training: 1 session with biofeedback (visual)

Outcome Measures: COP gait line, contact area, contact time, peak pressure, pressure time integral

Data Collection Timepoints: Baseline, while receiving biofeedback

COP gait line (mm): Medial shift from 0-80% of stance: 0-10% (MD=-1.4, ES=-0.3), 11-20% (MD=-1.4, ES=-0.3), 21-30% (MD=-1.8, ES=-0.4), 31-40% (MD=-2.0 ES=-0.5), 41-50% (MD=-2.2, ES=-0.5), 51-60% (MD=-2.4, ES=-0.5), 61-70% (MD=-2.6, ES=-0.5), 71-80% (MD=-2.4, ES=-0.5), 81-90% (MD=-1.4, ES=-0.3)

Contact area (*cm*²): Increased in medial midfoot (MD=2.1, ES=0.3) and hallux (MD=0.1, ES=0.1)

Peak pressure (kPa):

Decreased at lateral midfoot (MD=-10.8, ES=-0.6), central forefoot (MD=-51.9, ES=-1.2), and lateral forefoot (MD=-19.1, ES=-0.6), increased at hallux (MD=39.4, ES=0.7)

PTI (kPa*s):

Decreased at lateral heel (MD=-7.4, ES=-0.5) and lateral midfoot (MD=-6.8, ES=-0.5), increased at hallux (MD=18.6, ES=0.7)

Torp et al.

(7 control, 11 biofeedback)

18 CAI

Design: Randomized controlled trial

Gait Training: 8 sessions with biofeedback (auditory)

Outcome Measures: COP gait line, peak pressure, maximum force

Data Collection Timepoints: Baseline, 24-48 hours after last gait training session, 1-week after last gait training session

COP gait line (mm):

Immediately post:

Medial shift from 41-100% of stance: 41-50% (MD=-4.9, ES=-1.5), 51-60% (MD=-6.5, ES=-1.7), 61-70% (MD=-8.2, ES=-1.9), 71-80% (MD=-9.6, ES=-2.1), 81-90% (MD=-9.8, ES=-2.1), 91-100% (MD=-8.8, ES=-1.6)

1-week post: Medial shift from 31-50% and at 81-90% of stance: 31-40% (MD=-4.0, ES=-1.5), 41-50% (MD=-5.2, ES=-1.6), 81-90% (MD=-7.8, ES=-1.7)

Peak pressure (kPa):

Immediately post:

Decrease in lateral midfoot (MD=-22.2, ES=-1.3) and lateral forefoot (MD=-28.1, ES=-0.9), increased at medial forefoot (MD=36.0, ES=0.9)

1-week post: Decrease in lateral midfoot (MD=-20.0, ES=-1.1) and lateral forefoot (MD=-16.4, ES=-0.4)

Maximum Force (N):

Immediately post: Reduced in lateral midfoot (MD=-6.0, ES=-1.1) and lateral forefoot (MD=-6.8, ES=-1.5), increased in medial forefoot (MD=7.1, ES=1.47)

1-week post: Reduced in lateral midfoot (MD=-5.4, ES=-1.0) and lateral forefoot (MD=-4.3, ES=-1.1), increased in medial forefoot (MD=4.9, ES=1.0)

Abbreviations: CAI = chronic ankle instability, COP = center of pressure, ES = effect size, IC = initial contact, JCF = joint contact force, MD = mean difference, vGRF = vertical ground reaction force, 3D = three-dimensional.



Authors	Participants	Study Information	Main Findings
Donovan et al.	26 CAI	Design: Randomized controlled trial	Ankle Joint Angles (°): Increased dorsiflexion
	(13 control, 13 intervention)	<i>Gait Training:</i> 12 sessions with device (destabilization boot and sandal)	(MD=5.4, ES=3.4) during mid-late stance Knee Joint Angles (°):
		<i>Outcome Measures:</i> 3D joint angles for ankle, knee, hip for 1-100% of stride cycle	No significant differences Hip Joint Angles (°):
		<i>Data Collection Timepoints:</i> Baseline, 2-7 days after last gait training session	No significant differences
Koldenhoven et al.	27 CAI (14 control,	Design: Randomized controlled trial	Ankle Joint Angles (°): Decreased ankle inversion at IC (MD=-7.3, ES=-1.6)
	13 intervention)	Gait Training: 8 sessions with biofeedback (visual)	and throughout entire stride cycle (MD=-5.9, ES=-1.2)
		Outcome Measures: 3D joint angles for ankle, knee, hip	Vaca Isint Angles (9).
		Data Collection Timenoints: Baseline 24-72 hours after	Increased external rotation (MD=3.2, ES=0.7) during terminal swing
		last gait training session	
			Hip Joint Angles (°): No significant differences
Migel et al.	19 CAI	Design: Descriptive laboratory study with repeated measures	Ankle Joint Angles (°): Laboratory
		<i>Gait Training:</i> 2 sessions (1 laboratory, 1 real world) with biofeedback (haptic)	Increased abduction (MD=-1.7, ES=-1.0) during loading response
		<i>Outcome Measures:</i> 3D ankle, hindfoot, and forefoot joint angles during stance	<i>Real World</i> Decreased inversion (MD=-2.5, ES=-0.3) and increased abduction (MD=2.3, ES=0.5) during loading
		Data Collection Timepoints: Baseline, immediately after	response

Table 2. Summary of articles related to kinematic outcome measures. All results are reported in comparison to baseline values.



Participants	Study Information	Main Findings
15 CAI	Design: Randomized crossover laboratory study	RMS Amplitude Pre-IC Boot:
	<i>Gait Training:</i> 1 session with device (destabilization boot and sandal)	Increased for fibularis longus (MD=0.10, ES=0.9)
	<i>Outcome Measures:</i> RMS amplitude normalized to MVIC for 100ms pre-IC and 200ms post-IC for tibialis anterior, fibularis longus, lateral gastrocnemius, rectus femoris, and gluteus medius	Sandal: Increased for fibularis longus (MD=0.06, ES=0.7) RMS Amplitude Post-IC Boot:
	Data Collection Timepoints: Baseline, 2-7 days after	Increased for fibularis longus (MD=0.23, ES=1.3)
	last gait training session	<i>Sandal</i> Increased for fibularis longus (MD=0.14, ES=1.0)
26 CAI (13 control, 13 intervention)	 Design: Randomized controlled trial Gait Training: 12 sessions with device (destabilization boot and sandal) Outcome Measures: RMS amplitude normalized to quiet standing for 1-100% of stride cycle for tibialis anterior, fibularis longus, fibularis brevis, and medial gastrocnemius 	RMS Amplitude 1-100% gait cycle Decreased for fibularis longus during early stance (MD=2.9, ES=4.8) and mid-swing (MD=1.0, ES=2.5) phases of gait
	<i>Data Collection Timepoints:</i> Baseline, 2-7 days after last gait training session	
	Participants 15 CAI 26 CAI (13 control, 13 intervention)	ParticipantsStudy Information15 CAIDesign: Randomized crossover laboratory studyGait Training:1 session with device (destabilization boot and sandal)Outcome Measures:RMS amplitude normalized to MVIC for 100ms pre-IC and 200ms post-IC for tibialis anterior, fibularis longus, lateral gastrocnemius, rectus femoris, and gluteus mediusData Collection Timepoints:Baseline, 2-7 days after last gait training session26 CAI (13 control, 13Design: Randomized controlled triat (destabilization boot and sandal)0Outcome Measures: RMS amplitude normalized to quiet standing for 1-100% of stride cycle for tibialis anterior, fibularis longus, fibularis brevis, and medial gastrocnemiusData Collection Timepoints: Baseline, 2-7 days after last gait training session

Table 3. Summary of articles related to electromyography outcome measures. All results are reported in comparison to baseline values.

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Donovan et al.	10 CAI

Design: Descriptive laboratory study

Gait Training: 1 session with biofeedback (auditory)

Outcome Measures: RMS amplitude (not normalized due to within session testing design) for 200ms pre-IC and 200ms post-IC for tibialis anterior, fibularis longus, medial gastrocnemius, and gluteus medius

Data Collection Timepoints: Baseline, while receiving biofeedback

Feger & Hertel 10 CAI

Design: Descriptive laboratory study

Gait Training: 1 session with device (novel gait trainer with resistance bands)

Outcome Measures: RMS amplitude (not normalized due to within session testing design) for 200ms pre-IC and 200ms post-IC for tibialis anterior, fibularis longus, medial gastrocnemius, and gluteus medius.

Data Collection Timepoints: Baseline, while using device

RMS Amplitude Pre-IC Increased for fibularis longus (MD=80.2, ES=1.0)

RMS Amplitude Post-IC Increased for fibularis longus (MD=129.1, ES=0.8)

RMS Amplitude Pre-IC

No significant differences

RMS Amplitude Post-IC

Increased for fibularis longus (MD=200.1, ES=0.8) and medial gastrocnemius (MD=233.3, ES=0.7)

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Feger et al.	16 CAI	Design: Quasi-experimental trial	RMS Amplitude 0-100% of Stance
		<i>Gait Training:</i> 5 sessions with device (novel gait trainer with resistance bands)	Increased for fibularis longus from 21-60% and 82- 90% of stance phase: 21-30% (MD=2.4, ES=0.8), 31-40% (MD=2.2, ES=0.7), 41-50% (MD=3.1,
		<i>Outcome Measures:</i> RMS amplitude normalized to quiet standing for 0-100% of stance phase for tibialis anterior, fibularis longus, medial gastrocnemius, and gluteus medius.	ES=0.9), 51-60% (MD=2.8, ES=0.6), 81-90% (MD=2.1, ES=0.4) Decrease for gluteus medius 71-100% of stance
		Data Collection Timepoints:	phase: 71-80% (MD=-0.9, ES=-0.7), 81-90% (MD=- 1.0, ES=-0.9), 91-100% (MD=-1.6, ES=-0.9)
Knuckles et al.	12 CAI	Design: Descriptive laboratory study	RMS Amplitude Pre-IC
		<i>Gait Training:</i> 1 session with device (multi-axis destabilization device)	Wearing device: Increased for tibialis anterior (MD=3.6, ES=0.9)
		Outcome Measures: RMS amplitude normalized to quiet standing 50ms pre-IC and 200ms post-IC for	No significant differences
		tibialis anterior, fibularis longus, soleus, gluteus medius	RMS Amplitude Post-IC No significant differences
		<i>Data Collection Timepoints:</i> Baseline, while wearing device, immediately after devices removed	Post gait training: No significant differences







Figure 2. Meta-analysis results for the center of pressure (COP) gait line. Positive effect sizes indicate a lateral shift in COP. Negative effect sizes indicate a medial shift in COP. (a) indicates single session of gait training and data collected while receiving gait training. (b) indicates single session of gait training and data collected after gait training. (c) indicates multiple sessions of gait training, data collected 24-72 hours after gait training. (d) indicates multiple sessions of gait training, data collected 1-week after gait training.

A.) 0-10% of stance



C. 20-30% of stance



Effect Size

E. 40-50% of stance



Effect Size

G. 60-70% of stance



Effect Size

I. 80-90% of stance



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Figure 3. Meta-analysis results for the plantar pressure outcome measures. Positive effect sizes indicate an increase in pressure. Negative effect sizes indicate a decrease in pressure. (a) indicates single session of gait training and data collected while receiving gait training. (b) indicates single session of gait training and data collected after gait training. (c) indicates multiple sessions of gait training, data collected 24-72 hours after gait training. (d) indicates multiple sessions of gait training, data collected 1-week after gait training.



A.) Contact area central forefoot

C.) Peak pressure lateral forefoot



F.) Pressure time integral lateral midfoot



Figure 4. Meta-analysis results for the EMG outcome measures. Positive effect sizes indicate an increase in EMG activity. Negative effect sizes indicate a decrease in EMG activity. (a) indicates single session of gait training and data collected while receiving gait training.



A.) RMS amplitude post-IC for fibularis longus

Supplemental. Meta-analysis results for the non-significant findings for kinetic, kinematic, and electromyography outcome measures. (a) indicates single session of gait training and data collected while receiving gait training. (b) indicates single session of gait training and data collected after gait training. (c) indicates multiple sessions of gait training, data collected 24-72 hours after gait training. (d) indicates multiple sessions of gait training, data collected 1-week after gait training.

Kinetics

A.) Contact area - Lateral heel



D.) Contact area - Medial midfoot





J.) Contact time – Lateral heel



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M.) Contact time - Medial midfoot



P.) Contact time - Medial forefoot



S.) Contact time – Total foot



V.) Peak Pressure - Medial midfoot



Y.) Peak Pressure - Toes 2-5









HH.) Time to Peak Pressure – Medial heel





Kinematics

A.) Ankle Frontal Plane - Initial Contact



D.) Ankle Sagittal Plane - Loading Phase



Electromyography

A.) RMS Amplitude Pre-IC – Tibialis Anterior





D.) RMS Amplitude Post-IC - Tibialis Anterior