

1 **Title:** Is there a pathological gait associated with common soft tissue running injuries?

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16 **Title: Is there a pathological gait associated with common soft tissue running**
17 **injuries?**

18 **ABSTRACT**

19 **Background:** Previous research has demonstrated clear associations between specific
20 running injuries and patterns of lower limb kinematics. However, there has been minimal
21 research investigating whether the same kinematic patterns could underlie multiple
22 different soft tissue running injuries. If they do, such kinematic patterns could be considered
23 global contributors to running injury.

24 **Hypothesis:** Injured runners will demonstrate differences in running kinematics when
25 compared to injury free controls. These kinematic patterns will be consistent amongst injury
26 subgroups.

27 **Study Design:** Case- Control Study

28 **Methods:** We studied 72 injured runners and 36 healthy controls. The injured group
29 contained four subgroups of runners with either patellofemoral pain, iliotibial band
30 syndrome, medial tibial stress syndrome or Achilles tendinopathy (n = 18 each). Three-
31 dimensional running kinematics were compared between injured and healthy runners and
32 then between the four injured subgroups. A logistic regression model was used to
33 determine which parameters could be used to identify injured runners.

34 **Results:** The injured runners demonstrated greater contralateral pelvic drop and forward
35 trunk lean at mid-stance and a more extended knee and dorsiflexed ankle at initial contact.
36 The subgroup ANOVA found these kinematic patterns were consistent across each of the

37 four injury subgroups. Contralateral pelvic drop was found to be the most important
38 variable predicting classification of participants as healthy/injured. Importantly, for every 1°
39 increase in pelvic drop there was an 80% increase in the odds of being classified injured.

40 **Conclusion:** This study identified a number of global kinematic contributors to common
41 running injuries. In particular, we found injured runners to run with greater peak
42 contralateral pelvic drop and trunk forward lean, as well as an extended knee and
43 dorsiflexed ankle at initial contact. Contralateral pelvic drop appears to be the variable most
44 strongly associated with common running related injuries.

45 **Clinical Relevance:** The identified kinematic patterns may prove beneficial for clinicians
46 when assessing for biomechanical contributors to running injuries.

47 **Keywords:** Running, kinematics, injury, gait

48

49 **What is currently known about the subject:**

- 50 • Previous research has demonstrated clear associations between specific running
51 injuries and patterns of lower limb kinematics.
- 52 • Studies have found similar kinematic patterns that could underlie multiple different
53 running injuries.
- 54 • There may be kinematic patterns that represent global kinematic contributors to
55 running injury.

56 **What this study adds to existing knowledge:**

- 57 • The characteristics of increased contralateral pelvic drop, forward trunk lean and a
58 more extended knee and dorsiflexed ankle at initial contact are associated with
59 multiple common soft tissue running injuries.
- 60 • Contralateral pelvic drop was identified as the parameter most strongly associated
61 with running injury.
- 62 • For every 1° increase in contralateral pelvic drop there was an 80% increase in the
63 odds of being classified injured.
- 64 • This is the first kinematic study to identify a potential set of global kinematic
65 contributors to running injury.

66

67 **INTRODUCTION**

68 Running is an increasingly popular method of physical activity, however it also poses a risk
69 of injury to the musculoskeletal system. It has been reported that approximately 50% of
70 runners become injured annually with 25% injured at any one time.¹³ The majority of
71 running related injuries are considered to be overuse injuries, with the most frequently
72 injured sites including the knee, foot and lower leg, with incidence rates reported of around
73 50%, 39% and 32% respectively⁴⁶. Less common injury sites include the ankle and lower
74 back, as well as the hip and pelvis, with incidence rates ranging from 4% to 16%, 5% to 19%
75 and 3 to 11% respectively⁴⁵. Of all running related injuries, the most frequently cited injuries
76 include patellofemoral pain, iliotibial band syndrome, medial tibial stress syndrome, Achilles
77 tendinopathy, plantar fasciitis, stress fractures and muscle strains.^{24, 44} Many of these
78 injuries are known to have high reoccurrence rates, leading to a reduction or cessation of
79 training in approximately 30 to 90% of cases.⁴⁷ The factors related to the development of

80 running related injuries are multifactorial and diverse, however it is widely accepted that
81 abnormal running kinematics play a role.^{1, 7, 31}

82 There has been a large amount of research that has sought to identify the kinematic
83 patterns associated with many common soft tissue running injuries, including medial tibial
84 stress syndrome (MTSS)²⁶, patellofemoral pain (PFP)^{32, 52}, iliotibial band syndrome (ITBS)^{31 12}
85 and Achilles tendinopathy (AT)³⁹. Interestingly, many of these studies have reported similar
86 kinematic patterns to be associated with different running injuries. For example, increased
87 hip adduction has been associated with PFP^{32, 52} and ITBS^{31 12} and increased hip internal
88 rotation has been associated with PFP⁴¹ and MTSS²⁶. Research has also suggested that due
89 to the kinematic coupling between the femur, knee and foot, increased hip adduction or hip
90 internal rotation may contribute to greater rearfoot eversion ^{2, 27, 38}. Interestingly increased
91 rearfoot eversion has been associated with injuries such as MTSS ^{3, 50} and Achilles
92 tendinopathy.³⁹ This research suggests that there may be a number of similar kinematic
93 patterns that could underlie multiple different soft tissue running injuries. It is possible that
94 these patterns could lead to elevated stress on multiple anatomical structures leading to
95 injury development at different areas. These kinematic patterns may represent global
96 contributors to injury.

97 Recent research supports the idea of biomechanical parameters that could be considered
98 global contributors to running injury. In a prospective study of 249 runners, Davis *et al*⁷
99 reported that runners who went on to develop a range of different injuries, demonstrated
100 significantly elevated vertical loading rates. While in a retrospective study which
101 investigated runners with AT and MTSS, Becker *et al*³ reported greater rearfoot eversion at
102 late stance phase, to be a characteristic consistently associated with injury. Although these

103 two studies provide preliminary evidence for the existence of global contributors to running
104 injury, Davis *et al*⁷ did not include kinematic data, while Becker *et al*³ investigated only
105 MTSS and AT. Therefore, further research is required to understand whether there are
106 similar kinematic patterns that may underlie multiple different running injuries. This
107 understanding would be invaluable to clinicians as it could be used as a basis for both
108 screening techniques as well as preventative and rehabilitative programs.

109 The aim of this current study was to identify whether there are kinematic parameters that
110 may represent global kinematic contributors to running injury. To achieve this objective, we
111 sought to identify whether there are differences in running kinematics between a large
112 group of runners with common running injuries (ITBS, PFP, MTSS and AT) compared to a
113 healthy control group. We hypothesised that the pooled group of injured runners would
114 demonstrate greater contralateral pelvic drop, hip adduction and rearfoot eversion angles
115 when compared to injury free controls. We also hypothesised that these kinematic patterns
116 would be consistent amongst injury subgroups.

117 **METHODS**

118 **Participants**

119 A total of 108 runners were enrolled in this current study, including 72 injured runners (28
120 males, 44 females) and 36 healthy controls (15 males, 21 females) matched for age, height
121 and weight (Table 1). The injured group contained subgroups of 18 runners with PFP, ITBS,
122 MTSS and AT (Table 2). These injuries were selected as they are cited as the most prevalent
123 soft tissue overuse running injuries.²⁴ An a priori sample size calculation was conducted
124 using data from a previous study reporting kinematic differences between healthy and
125 injured runners.³² Using g*power software, we calculated that we would need at least 98

126 people (65 injured) in order to detect an effect size of 0.75 with a power of 0.85 and a
 127 critical $\alpha = 0.01$. Participants were recruited via poster advertisements at local running clubs
 128 and sports injury clinics. All participants provided written informed consent prior to
 129 participation and ethical approval was obtained via the local ethics committee.

130

	Healthy (n = 36)	Injured (n = 72)
Age (years)	33.2 (8.4)	34.8 (9.9)
Mass (kg)	60.8 (8.4)	63.4 (10.5)
Height (cm)	171.6 (7.3)	170.7 (8.6)
BMI (kg.m ⁻²)	20.6 (1.8)	21.7 (2.7)
Miles run per week*	60.5 (23.2)*	21.2 (13.1)*

131 **Table 1: Mean (SD) participant characteristics. *indicates statistical significance at $p < 0.01$.**

	PFP (n = 18)	ITBS (n = 18)	MTSS (n = 18)	AT (n = 18)
Age (years)	34.5 (9.4)	34.3 (7.9)	31.9 (9.7)	38.5 (11.7)
Mass (kg)	64.4 (9.6)	63.6 (11.2)	62.5 (10.1)	63.1 (11.8)
Height (cm)	173.5 (8.5)	170.6 (8.5)	167.3 (8.1)	171.6 (8.7)

BMI (kg.m ⁻²)	21.3 (1.9)	21.8 (3.3)	22.2 (2.3)	21.3 (2.0)
Miles run per week*	18.6 (6.9)	14.8 (5.8)	19.5 (12.2)	31.9 (17.6)*

132 **Table 2: Mean (SD) injury subgroup characteristics. *indicates statistical significance at p = <0.01.**

133 **Inclusion/ Exclusion Criteria**

134 **Injured Group**

135 The injured group included individuals with a current diagnosis of either PFP, ITBS, MTSS or
136 Achilles tendinopathy. Injury diagnosis was confirmed following a physical examination by a
137 qualified physiotherapist in accordance with previously published diagnostic criteria for
138 PFP,⁶ ITBS,¹⁷ MTSS⁵⁴ and Achilles tendinopathy²² (Supplementary File 1). All participants
139 reported being able to run up to 10 minutes before the onset of pain and maximal pain
140 during running greater than 3/10 on a numerical rating scale (0 = no pain, 10 = worst
141 possible pain). Additionally, all participants reported they were not currently receiving
142 medical treatment for their injury and that their pain had caused a restriction to their
143 running volume and/or frequency for a minimum of 3 months. Previous research has
144 reported training factors such as increases in weekly training volume, to increase the risk of
145 injury. This is likely due to a sudden excessive rise in acute tissue stress on the
146 musculoskeletal system, resulting in insufficient time for adaptive changes³³. Therefore, in
147 order to control for training errors as a cause of injury, participants were excluded if they
148 reported an increase in weekly training volume of greater than 30% preceding the onset of
149 injury.

150 **Control Group**

151 Control participants were included if they reported running a minimum of 30 miles per week
152 for the last 18 months with no reported injury. Participants were excluded if they reported
153 any musculoskeletal ailment within the last 18 months that caused a restriction or cessation
154 of running, or any need to seek advice from a health care professional. Additional exclusion
155 criteria included previous history of overuse running injury, injury caused by another sport,
156 previous spinal injury or lower limb surgery.

157 **Procedures**

158 Kinematic data were collected from all participants whilst running on a treadmill at 3.2m/s
159 wearing their own running shoes. After a 5 minute warm up period, 30 seconds of kinematic
160 data were collected using a 12 camera Qualysis Oqus system (240Hz). A total of nine
161 anatomical segments were tracked following a previously published protocol by the same
162 authors shown to have good to excellent repeatability.^{28, 37} Segments included the thorax,
163 pelvis and bilateral thigh, shank and foot segments. In addition, a further rearfoot segment
164 was included using 3 non colinear markers attached to the heel of the participant's shoes.
165 The foot segment was used to calculate sagittal plane ankle kinematics while the rearfoot
166 segment was used to calculate frontal plane foot kinematics. Further details of the markers
167 used to track each segment and the precise definition of the anatomical coordinate systems
168 is provided in supplementary file 2 and described in previous publications.^{14, 28, 37}

169 Raw kinematic data were low pass filtered at 10Hz. Intersegmental kinematics, along with
170 the motions of the pelvis and thorax with respect to the laboratory system, were calculated
171 using a six degrees of freedom model using the commercial software Visual 3D (C-Motion).
172 Gait events were defined using a kinematic approach²⁰ and subsequently used to segment
173 each kinematic signal into a minimum of 10 consecutive gait cycles. An ensemble average

174 for each signal was created and selected kinematic parameters derived from the ensemble
175 average curves. This latter processing was carried out using a custom Matlab script.

176 **Data Analysis**

177 A range of kinematic parameters at both initial contact and mid-stance were selected for
178 analysis. Parameters at initial contact included sagittal plane angles of the trunk, pelvis, hip,
179 knee and ankle as well as frontal plane angles of the trunk and rearfoot. Peak angles at mid
180 stance included sagittal and frontal plane angles of the trunk, pelvis, knee and ankle and
181 rearfoot as well as transverse plane angles of the hip and knee. Parameters were selected
182 based on previous research reporting differences between injured and non-injured
183 runners^{39, 41, 52}. Peak angles at mid-stance were defined as the maximum joint angle
184 between initial contact and toe off. Foot strike patterns of each group were determined
185 based on the kinematic waveforms of the ankle joint. Where the ankle demonstrated an
186 immediate movement into plantarflexion, participants were classified as having a rearfoot
187 strike, participants demonstrating immediate ankle dorsiflexion were classified as a forefoot
188 strike. The injured leg was analysed from the injured runners, right or left leg was analysed
189 at random from the healthy runners in order to match the total distribution of right and left
190 legs in the injured group.

191 **Statistical Analysis**

192 Participant characteristics were analysed using independent t-tests for the healthy versus
193 injured group comparisons and a one-way ANOVA for the subgroup analysis (Table 1 & 2).
194 Chi-squared tests were used to assess for differences in distribution of foot strike patterns
195 between the groups. In order to identify possible global contributors to running injury we
196 used a two-phased approach. Firstly, data from the injured group were pooled and

197 kinematic parameters compared with those of the healthy group using an independent t-
198 test. Secondly, for any variables found to be significant different following the injured versus
199 healthy comparison, we assessed for subgroup differences between the four injury
200 subgroups using a one-way ANOVA test and post hoc Least Significant Difference (LSD). In
201 order to be considered a global contributor to running injury, we required a kinematic
202 parameter to be consistent across the different injury groups. This ensured that differences
203 observed in the pooled injury data, were not the result of large effects in one of the injury
204 subgroups. Before analysis, all kinematic parameters were assessed for homogeneity of
205 variance and normal distribution using Levine's test ($p = >0.05$) and Shapiro-Wilk ($p = >0.05$).
206 Where assumptions were not met, an equivalent non-parametric test was used. In order to
207 reduce the possibility of type I error, a critical $\alpha = 0.01$ was used for injured versus healthy
208 comparisons. However, we used a critical $\alpha = 0.05$ for the subgroup ANOVA analysis, due to
209 the smaller subgroup sample sizes. This was deemed appropriate given the smaller number
210 of group comparisons and therefore lower likelihood of type I error.

211 In addition to calculating statistical significance for group comparisons, we also calculated
212 effect sizes. For t-test comparisons, we used Cohen's D and interpreted an effect size of 0.2,
213 0.5 and 0.8 as small, medium and large respectively.⁴ For the ANOVA comparisons, we used
214 the Eta squared statistic ($\eta^2 = SS \text{ between groups} / SS \text{ total}$) and interpreted effect sizes of
215 0.01, 0.09 and 0.25 as small, medium and large respectively.⁴

216 Finally, a forward stepwise binary logistic regression analysis was conducted in order to
217 determine which kinematic parameters could predict classification into either the injured or
218 the healthy group. Parameters identified to be significantly different between healthy and

219 injured groups were considered for the regression model. Variables were excluded from the
220 regression model if they were found to demonstrate differences between injury subgroups.

221

222 **RESULTS**

223 **Injured versus Healthy**

224 The pooled data showed the injured runners to land with significantly more knee extension
225 and ankle dorsiflexion (Table 3, Figure 2). At mid-stance, the injured runners were found to
226 have significantly greater forward trunk lean, CPD (Figure 1a) and hip adduction (Figure 1c &
227 3, Table 4). Large effect sizes of 1.37, 0.89 and 0.87 were observed for CPD, hip adduction
228 and knee flexion at initial contact respectively (Table 3 & 4). Trunk forward lean at mid-
229 stance and ankle dorsiflexion at initial contact demonstrated moderate effect sizes of 0.65
230 and 0.71 respectively (Table 3 & 4). Chi-squared tests found no significant difference in the
231 distribution of foot strike patterns between the groups ($p = 0.332$). In the healthy group
232 there was a total of 17 forefoot and 19 rearfoot runners. In the Injured group there was a
233 total of 27 forefoot and 45 rearfoot runners.

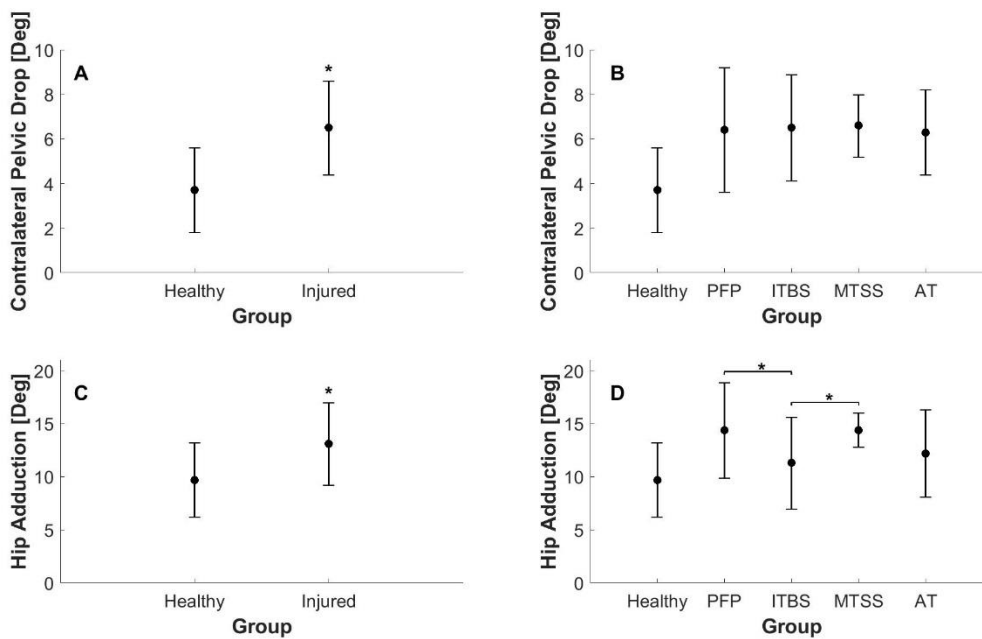
234

Variable	Control	Injured	P-value	Effect Size
Trunk Forward Lean (°)	3.9 (2.9)	5.7 (3.9)	0.033	0.52
Trunk Ipsilateral Lean (°)	2.5 (1.8)	3.1 (2.2)	0.257	0.28

Pelvis Anterior Tilt (°)	5.9 (3.3)	7.0 (3.8)	0.132	0.32
Knee Flexion* (°)	10.2 (4.8)	6.0 (4.9)	<0.01*	0.87
Ankle Dorsiflexion* (°)	2.4 (6.5)	7.2 (6.9)	<0.01*	0.71
Rearfoot Inversion (°)	8.7 (6.1)	6.2 (4.5)	0.018	0.47

235 **Table 3: Kinematic parameters at initial contact. Data represents angle at initial contact in degrees. ***
 236 **indicates statistical significance at $p < 0.01$.**

237



238

239 **Figure 1: A: Contralateral pelvic drop for healthy and injured groups. B: Contralateral pelvic drop for healthy**
 240 **and injury subgroups. C: Hip adduction for healthy and injured groups. D: Hip adduction for healthy and**
 241 **injury subgroups. PFP = patellofemoral pain, ITBS = iliotibial band syndrome, MTSS = medial tibial stress**
 242 **syndrome, AT = Achilles tendinopathy. Whiskers represent +/- 1SD. * indicates statistically significant**

243 differences for T-Tests (A & C) and subgroup ANOVA (B & D). Healthy group is shown in B & D for

244 comparison purposes only.

245

Variable	Control	Injured	P value	Effect Size
Trunk Forward Lean* (°)	9.5 (2.9)	12.0 (4.9)	<0.01*	0.65
Trunk Ipsilateral Lean (°)	3.6 (1.8)	4.3 (2.6)	0.094	0.33
Pelvis Anterior Tilt (°)	5.0 (2.9)	5.7 (3.8)	0.553	0.19
Contralateral pelvic drop* (°)	3.7 (1.9)	6.4 (2.1)	<0.01*	1.37
Hip Adduction* (°)	9.7 (3.5)	13.0 (3.9)	<0.01*	0.89
Hip internal rotation (°)	4.4 (6.8)	4.2 (8.0)	0.874	0.03
Knee Flexion (°)	32.7 (4.9)	32.3 (5.0)	0.556	0.09
Knee Adduction (°)	-1.9 (3.1)	-2.0 (3.5)	0.785	0.06

Knee External Rotation (°)	6.7 (5.5)	7.1 (6.9)	0.616	0.06
Ankle Dorsiflexion (°)	22.3 (2.9)	21.9 (4.3)	0.964	0.09
Rearfoot Eversion (°)	2.6 (3.2)	4.0 (3.5)	0.047	0.42

246 **Table 4: Peak kinematic angles during stance phase. Data represents maximum joint angle between initial**
247 **contact and toe off. * indicates statistical significance at p <0.01.**

248 **Injury Subgroups**

249 The subgroup ANOVA analysis was conducted in order to identify if there were differences
250 between injury subgroups for variables identified as being different between the pooled
251 injured and healthy groups. This analysis found no differences for ankle dorsiflexion and
252 knee flexion at initial contact (Table 5). Furthermore, there were no differences in peak
253 trunk forward lean and CPD during mid-stance (Table 5), indicating these parameters were
254 consistent across the injury subgroups. However there was a significant difference between
255 injury subgroups for peak hip adduction (Table 5). Post hoc LSD tests found the PFP (p =
256 0.018) and MTSS (p = 0.016) groups to have 3.1° and 3.2° more hip adduction than the ITBS
257 group (Figure 1d).

258

	PFP	ITBS	MTSS	AT	ANOVA Between Injury Groups	Effect Size Eta Squared (η^2)
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Initial Contact						
Knee Flexion (°)	5.5 (4.6)	6.6 (5.7)	4.7 (5.2)	7.4 (4.1)	0.365	0.05
Ankle Dorsiflexion (°)	10.6 (3.9)	7.1 (5.6)	5.5 (9.2)	5.6 (7.1)	0.088	0.09
Mid Stance						
Trunk Forward Lean (°)	11.9 (5.1)	14.3 (5.5)	10.9 (4.9)	11.3 (3.4)	0.160	0.07
Contralateral Pelvic Drop (°)	6.4 (2.8)	6.5 (2.4)	6.6 (1.4)	6.3 (1.9)	0.986	0.002
Hip Adduction* (°)	14.4 (4.5)	11.3 (4.3)	14.4 (1.6)	12.2 (4.1)	0.032*	0.12

259 **Table 5: Between injury subgroups ANOVA. * indicates statistical significance at p <0.05**

260 **Logistic Regression**

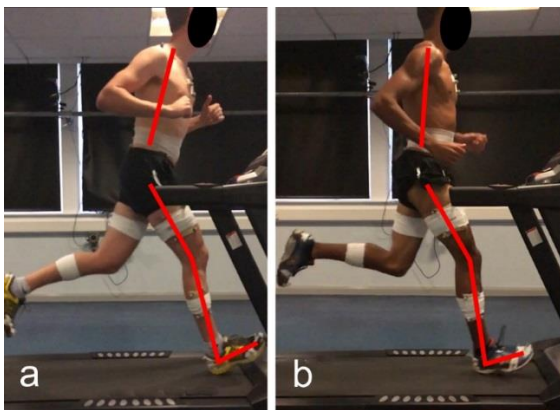
261 The final variables identified as global kinematic contributors included knee flexion and
262 ankle dorsiflexion at initial contact as well as trunk forward lean and CPD at mid-stance. All
263 four variables were entered into the logistic regression model. The forward stepwise logistic
264 regression model identified that CPD at mid-stance (OR = 1.87; 95% CI: 1.41, 2.49; p =
265 <0.001) and knee flexion at initial contact (OR = 0.87; 95% CI: 0.78, 0.97; p = 0.012) were
266 significant predictors of classification as either healthy or injured, explaining 47% of the
267 variance in the data ($R^2 = 0.466$). The most important predictor variable was CPD, with an

268 80% increase in the odds of being classified injured for every 1° increase in pelvic drop. For
269 knee flexion there was a 23% reduction in the odds of being classified injured for every 1°
270 increase in knee flexion at initial contact.

271

272 **DISCUSSION**

273 This study identified a number of kinematic differences between the injured and healthy
274 runners that were consistent across injury subgroups. In particular the injured runners were
275 found to demonstrate significantly greater peak contralateral pelvic drop (CPD) and forward
276 trunk lean, as well as a more extended knee and dorsiflexed ankle at initial contact (Table 3,
277 4 & 5) (Figures 2 & 3). We found CPD to be the most important predictor variable when
278 classifying runners as healthy or injured. These kinematic patterns may represent global
279 kinematic contributors to soft tissue running injuries and together may define a pathological
280 running gait.



281

282 **Figure 2: Two dimensional representation of forward trunk lean, knee flexion and ankle dorsiflexion angles at initial**
283 **contact. A = injured runner, B = healthy runner.**

284



285

286 **Figure 3: Two dimensional representation of contralateral pelvic drop and hip adduction during mid-stance. A = injured**
 287 **runner, B = healthy runner.**

288

289 **Global kinematic contributors**

290 Peak contralateral pelvic drop was found to be the kinematic parameter most strongly
 291 associated with running injury. Previous studies have associated CPD with PFP⁵² and MTSS,²⁶
 292 however this study identified increased CPD amongst multiple different running related
 293 injuries, including ITBS and Achilles tendinopathy (Figure 1b). Therefore, CPD may represent
 294 a global kinematic contributor and risk factor for many common soft tissue running injures.

295 It is likely that CPD will influence lower limb tissue stress at a number of different
 296 anatomical sites through a number of different mechanisms. For example, Tateuchi *et al*⁴³
 297 identified that increasing CPD resulted in an increase in iliotibial band tension at the lateral
 298 femoral condyle. This will likely influence ITBS development through increased strain rate¹⁹
 299 and increased compression between the ITB and lateral femoral condyle¹¹. At the same
 300 time, an increase in ITB tension will result in a lateral displacement of the patella.²⁹ Lateral
 301 displacement of the patella will lead to a rise in patellofemoral joint stress, leading to PFP
 302 development,³⁶ while at the lower limb, increased CPD will result in a medial shift in the
 303 ground reaction force relative to the knee joint centre.^{37, 42} This may alter the force

304 distribution through the lower limb, leading to increased bending forces on the medial tibia⁵
305 and potentially alter pressure distribution through the foot. This may contribute to the
306 development of either MTSS or AT.^{25, 50}

307

308 One possible explanation for the increased CPD observed in the injured group could be due
309 to reduced strength or neuromuscular function at the hip. Previous authors have reported
310 delayed onset of gluteus medius and maximus in runners with PFP⁵¹ and AT¹⁵, while others
311 have reported reduced hip abductor strength in runners with ITBS¹⁶, PFP⁴¹, AT¹⁸ and MTSS⁴⁸.
312 The hip abductors, in particular the gluteus medius, are thought to control frontal plane
313 kinematics of the pelvis and hip⁴⁰. Therefore, it is conceivable that reduced strength or
314 neuromuscular function of the gluteus medius would lead to an inability to stabilise the
315 pelvis in the frontal plane, causing increased CPD.

316

317 We also found the injured runners to land with greater knee extension and ankle
318 dorsiflexion (Table 3, Figure 2), which may influence tissue stress in a number of ways.
319 Firstly, in knee extension the patella becomes vulnerable to lateral tilt and displacement
320 which may influence patellofemoral contact areas and joint stress during early stance³⁵.
321 Secondly, an extended knee and dorsiflexed ankle at initial contact is typically associated
322 with a greater distance between the centre of mass and the foot at contact. Greater
323 distance between the centre of mass and foot, as well as larger ankle dorsiflexion angles,
324 have been associated with increased knee joint loading and braking impulse⁴⁹. An
325 extended knee at initial contact has also been reported to reduce the ability to attenuate
326 impact forces during early stance⁸. Collectively it seems plausible that the extended lower

327 limb posture at initial contact may influence impact loading and knee joint loading during
328 early stance.

329

330 One possible mechanism explaining the differences in forward trunk lean may be due to
331 strength deficits around the gluteals and paraspinals. Previous studies have reported fatigue
332 of the paraspinal and gluteal muscles to be associated with an increase in trunk forward
333 lean during running²¹ and drop landings²³. Therefore, reduced strength capacity of the
334 gluteals and paraspinals may result in an inability to maintain an upright running posture
335 amongst the injured runners.

336

337 **Kinematic Subgroups**

338 While hip adduction was found to be greater amongst the pooled injured group, the
339 subgroup analysis revealed this parameter differed across the injury subgroups (Table 5,
340 Figure 1c & 1d). Specifically, we found hip adduction to be greater amongst subgroups of
341 runners with PFP and MTSS compared to the ITBS subgroup (Figure 1d). This finding is in
342 contrast to previous studies by Noehren et al³¹ and Ferber et al¹² who reported increased
343 hip adduction amongst runners with ITBS. One potential reason for the contrasting findings
344 may be due to sex differences between studies. Hip adduction has been reported to be
345 influenced by sex subgroups⁵² with greater hip adduction amongst female runners. In the
346 current study we included a mix of males and females while Noehren et al³¹ and Ferber et
347 al¹² only included female participants. While we acknowledge that hip adduction may be
348 an important kinematic risk factor for certain injuries, we feel our data suggests hip

349 adduction may be more influential in specific subsets of runners and pathologies, rather
350 than others.

351

352 **Limitations**

353 One limitation is that the study was retrospective and therefore it is not possible to
354 conclude if the observed kinematic patterns are the cause of injury, or the result of injury.
355 Nevertheless, we ensured that all data were recorded before the onset of pain to minimise
356 any possible effect of pain on the observed kinematic patterns. However we cannot rule out
357 the possibility that participants may have adapted their running kinematics in response to
358 chronic injury or in apprehension of the acute onset of pain. Therefore, we acknowledge
359 that future prospective studies are required to further investigate whether the kinematic
360 patterns observed within the current study are the cause or effect of injury. Another study
361 limitation is the higher weekly mileage of the control group (Table 1). However, we feel that
362 this could be considered a strength, as previous research suggests running greater than 40
363 miles per week is a risk factor for developing injury.⁴⁶ On average, our healthy control group
364 were exceeding this threshold for more than 18 months prior to testing yet remained injury
365 free. Therefore, we feel the control group may be representative of a healthy running gait in
366 order to remain injury free at training loads exceeding the previously reported injury
367 threshold. It is also important to note that this study was limited to a select number of
368 common soft tissue running injuries and therefore these results may not apply to other
369 injuries such as plantar heel pain, stress fractures and muscle strains. Further research
370 would be required in order to establish a link between the identified kinematic patterns and
371 other running related injuries.

372 **Clinical Relevance**

373 The findings from the present study may have a number of clinical implications. Firstly, all of
374 the identified kinematic parameters can be easily visualised using two dimensional gait
375 analysis methods^{9, 10, 34} (Figures 2 & 3). A number of recent publications have shown 2D
376 assessments of CPD, hip adduction, trunk forward lean and sagittal plane knee and ankle
377 angles to be highly correlated with 3D measurement systems and to demonstrate high intra
378 and inter-tester reliability^{9, 10, 34}. Therefore, it should be possible to use 2D measurement
379 techniques to assess the biomechanical parameters which were associated with injury in
380 this study. Secondly, many of the identified global kinematic contributors to injury, can be
381 modified through gait retraining. For example, CPD and hip adduction angles can be
382 retrained using mirror feedback,⁵³ while knee and ankle angles are influenced by increasing
383 cadence or modifying foot strike patterns.³⁰ Therefore, this study highlights a number of key
384 kinematics that can be considered global contributors to running injury and can be easily
385 assessed and modified in clinical practice. This may assist clinicians in the development of
386 rehabilitation programs for common running related injuries.

387 **CONCLUSION**

388 This study identified a number of global kinematic contributors to common running injuries.
389 In particular, we found injured runners to run with greater peak contralateral pelvic drop
390 and trunk forward lean, as well as an extended knee and dorsiflexed ankle at initial contact.
391 Contralateral pelvic drop appears to be the variable most strongly associated with common
392 running related injuries. The kinematic patterns identified as global contributors to injury
393 can be easily assessed and modified in clinical practice.

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