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1 **Are the human leg joints controlled relative to a misaligned**
2 **standing reference?**

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16

17 **Abstract**

18 In human quiet standing, the relative position between ankle joint centre and line of gravity is
19 neurally regulated within tight limits. The regulation of the knee and hip configuration is
20 unclear and thought to be controlled passively. However, perturbed standing experiments
21 have shown a leg multi-joint coordination. Here, measuring the relative alignment between
22 leg joints and the line of gravity in quiet standing after walking, we investigated whether the
23 configuration is maintained over time through passive mechanisms or active control.

24

25 Thirteen healthy adults walked without following a path and then stood quietly for 7.6s on a
26 force platform (up to four trials). The transition between initiation and steady-state standing
27 (7.6s) was measured using motion capture. Sagittal leg joint centres' position relative to line
28 of gravity (CoG_{AP}) and their time constants were calculated in each trial. Ankle, knee and hip
29 joint moments were also calculated through inverse dynamics.

30

31 After walking, the body decelerated ($\tau=0.16s$). The ankle and hip joints' position relative to
32 CoG_{AP} measured at two time intervals of quiet standing (Mid=0.5-0.55s; End=7.55-7.6s)
33 were different (mean \pm sem CoG_{AP}-Ankle_Mid=47 \pm 4mm, CoG_{AP}-Ankle_End=58 \pm 5mm; CoG_{AP}-
34 Hip_Mid=2 \pm 5mm, CoG_{AP}-Hip_End=-5 \pm 5mm).

35 The ankle, knee and hip flexion-extension moments significantly changed.

36

37 Changes in joints position relative to CoG_{AP} and misalignment suggest that joint position is
38 not maintained over 7.6s, but regulated relative to a standing reference. Higher joint moments
39 at steady-state standing suggest mechanisms other than passive knee and hip regulation are
40 involved in standing.

41

42 **Keywords**

43 Human stance control, Standing balance, Neural control of movement, Leg joints, Body
44 misalignment to line of gravity, Initiation of standing

45 **Introduction**

46 Human standing is an everyday activity, and it constitutes the mechanical and control basis
47 for other movements, such as gait and reaching. In quiet standing, the body is unstable in the
48 sagittal plane (Morasso and Schieppati 1999; Loram et al., 2007; Kiemel et al., 2011). To
49 maintain standing successfully, the location of the body centre of mass and of the ankle joint
50 relative to the line of gravity need to be regulated via neural feedback control (Peterka and
51 Benolken 1995; Loram and Lakie 2002; Loram et al., 2007).

52

53 Because the body centre of mass sway range is only 18-21mm in standing (Gatev et al.,
54 1999), whole body configuration (i.e. the position of body segments and joints relative to the
55 line of gravity) must to be tightly regulated. The traditional understanding is that in quiet
56 standing, only the ankle position needs to be regulated. The knee and hip positions relative to
57 the line of gravity are thought to be either passively determined taking advantage of the close
58 packed position (Steindler, 1964, page 330-349; MacConaill & Basmajian, 1977, page 31-
59 52), or tonically but not phasically regulated (Steindler, 1964, page 106-108 and 110-114).
60 Essentially, it was accepted that there is no need for a modulation of knee and hip moments in
61 quiet standing.

62

63 More recent work has shown that ankle, knee, hip, L5-S1 joint (5th lumbar and 1st sacral
64 vertebrae), C7-T1 joint (7th cervical and 1st thoracic vertebrae) and atlanto-occipital joint are
65 controlled in a coordinated fashion in standing (Hsu et al., 2007) according to the
66 uncontrolled manifold analysis. Focusing on the hip joint, Kiemel and colleagues showed that
67 intrinsic stiffness is not enough for hip passive stability and neural control is required to
68 maintain standing (Kiemel et al., 2008). Furthermore, ankle, knee and hip joints showed a
69 multi-joint coordinated behaviour in perturbed standing (Di Giulio et al., 2013). When gentle
70 knee perturbations were applied at the knee, if the knee displacement after perturbation was
71 small, also ankle and hip displacements were reduced and the whole leg was stiffer (locked or
72 inverted-pendulum like). On the other hand, when the knee displacement after perturbation
73 was larger, ankle and hip displacements were also larger and the whole leg did not show an
74 inverted pendulum-like configuration. This suggests that leg joints' stiffness or mobilisation
75 is controlled collectively, and even the knee joint is not necessarily passively locked. What
76 remains an open questions is whether this inter-joint relationship is purely mechanical or
77 tonic or whether phasic control is involved.

78

79 We designed an experiment that substantially changed body configuration in order to
80 measure how joint position in relation to line of gravity was attained in the transition to quiet
81 standing. Gait before quiet standing was used to measure standing initiation and
82 configuration changes to achieve quasi-static equilibrium. We did not use non-ecological
83 perturbations, such as platform translations or tilts, to avoid the introduction of artificial
84 responses and habituation to the perturbation over time. Our approach was to study the
85 transition of joint position between initiation and steady-state standing and analyse which
86 factors could explain the process. By studying initiation of standing and transition to steady-
87 state standing, we asked (i) What is the relative alignment between leg joints and the line of
88 gravity? (ii) Is the leg configuration at steady state standing determined by the position at
89 initiation of standing or is it actively controlled and corrected?

90

91 One could expect that biomechanical (e.g. body deceleration to stop the body after walking)
92 and passive mechanisms (e.g. stiffness) could wholly explain the joint position at initiation
93 and steady-state standing. An additional expectation is that steady-state standing
94 configuration could depend on the body configuration at initiation of standing and no

95 corrections are occurring as long as standing is successful and efficient. Furthermore, if
96 optimisation and energy cost minimisation was a principle of standing regulation, steady-state
97 standing configuration should be consistent with reduced muscular effort. This would suggest
98 that steady-state standing configuration is aligned with the vertical to reduce load on the
99 joints and the need for phasic muscular activation to maintain balance.

100

101 On the other hand, if leg configuration changed during standing, we could investigate
102 whether the difference in leg configuration between initiation and steady-state standing was
103 consistent with energy cost minimisation (i.e. the joints became more aligned) or not. We
104 could also measure whether the steady-state leg configuration was dependant on the initial
105 variable condition established by gait (i.e. not repeatable and inconsistent across trials).

106

107 In this study, we measured the leg joint alignment with line of gravity in quiet standing and
108 we investigated the mechanisms involved in this task. Understanding whether alignment was
109 maintained or corrected would indicate whether passive stiffness or other mechanisms to
110 control the leg joints are involved in standing.

111 **Methods**

112 *Ethical approval*

113 Participants gave written informed consent to these experiments which were approved by the
114 ethics committee of the Institute for Biomedical Research into Human Movement and Health,
115 Manchester Metropolitan University and conformed to the standards of the Declaration of
116 Helsinki.

117
118 *Participants and procedures*

119 Thirteen participants (age 46 ± 13 years, mass 71.7 ± 13.0 kg, height 1.68 ± 0.13 m, seven
120 women and six men) who self-reported no neurological or musculoskeletal injuries or
121 disorders took part in this study.

122
123 The data reported here is part of a larger experiment that lasted about 3 hours. For each
124 participant, the session was structured as follows. Participants arrived to the laboratory and
125 informed consent was obtained (10 min). Bilateral knee and hip MRIs were collected (60
126 min) and markers were placed on participant's anatomical landmarks (30 min). The first two
127 trials of the current experiment (5 min) were collected and then an intervening knee
128 perturbation experiment (40 min including EMG placement, as reported in Di Giulio et al.
129 2013) was conducted. The remaining trials of the current experiment were recorded after that
130 (5 min), and a final experiment on control of standing with another set-up (30 min) was
131 recorded. Breaks were also allowed between trials and experiments.

132
133 For this experiment, participants walked randomly for a few seconds around the laboratory
134 without following a particular path and ended their walk anywhere on a force plate
135 (508x464mm) with feet broadly symmetrical (about shoulder width, feet broadly parallel to
136 each other). Some participants walked over an imaginary circle or ellipsoid, others walked on
137 a straight line, others turned around and changed direction at least once. Participants were
138 asked to approach the force platform in a straight line (last 1-2 steps). The operator monitored
139 the participant's gait phase in order to start the recording timely, and the trial was repeated if
140 the participants did not approach the platform complying with this criterion. Participants were
141 not asked to replicate their walking path and most of them completely changed it over
142 different trials. Participants were asked to end their walking phase in a comfortable and usual
143 manner, and stand normally looking in front of them.

144
145 The recording was manually initiated when the participant approached the force plate and
146 each trial lasted 30s (from when the participant approached the force plate). Because of this
147 variable trial start, different effective standing durations were recorded. For analysis, the
148 longest common duration of standing after its initiation (flat feet time, see below) was used
149 (7.6s, see Fig. 1). Although a longer common duration was not possible, 7.6s after standing
150 initiation is likely to be sufficient to measure changes due to and possibly beyond body
151 deceleration, without fatiguing the participants. All the participants performed at least four
152 trials (with intervening break). When technical problems were identified in real time, the trial
153 was repeated. However, other technical difficulties in the markers trajectories reconstruction
154 were only identified during data processing and those trials were not included in further
155 analysis. For this reason, out of the thirteen participants, we could use 4 trials for seven
156 participants, 3 trials for three participants and 2 trials for three participants. These
157 participants were included in the analysis because more than one repetition was available, to
158 include as much population variability as possible, and because the data included was highly
159 reliable and accurate thanks to the precision of the techniques (motion capture and force

160 plates) and the corrections adopted from the MRI scans (marker positioning and correction of
161 joint centres calculation).

162

163 ***Apparatus and measurements***

164 *Imaging*

165 Four MRI scans were collected with the participants in the standing position (G-Scan, Esaote,
166 Genoa, Italy) to improve joint location accuracy. The same protocol was used for knee and
167 hip joints bilaterally: Spin T1-weighted HF, matrix 256x256, coronal and transverse planes.
168 Slice thickness and the inter-slice gap were 0.4mm and 4mm for the knees, and 0.6mm and
169 6mm for the hips. Cod liver oil pills were placed on anatomical landmarks where the retro-
170 reflective markers would be placed for motion analysis. If the image showed that the cod
171 liver oil pill was not placed correctly, it was replaced accordingly and another set of scans
172 was collected. This accurate location was then used to place the motion analysis marker.

173

174 *Motion capture*

175 A ten camera motion analysis system (VICON 612, Oxford Metrics, UK) was used to
176 measure body kinematics. Retro-reflective markers were placed on the sacrum, third lumbar
177 vertebral process (L3), twelfth, tenth, seventh and third thoracic vertebral process (T12, T10,
178 T7, T3), seventh cervical vertebral process (C7) and sternum and clavicle. Other markers
179 were placed bilaterally on the first, second and fifth metatarsal head, the lateral and medial
180 malleolus, the heel, the tibia (for 3D segment definition), and the most prominent points of
181 the lateral and medial tibial condyles, the lateral and medial femoral epicondyle, the greater
182 trochanters, the anterior and posterior iliac spines, the zygomatic process anterior to the
183 auditory meatus, and the temporal process of the zygomatic bone (at the inferior margin of
184 the ocular orbit). After walking, participants stood with both feet on a force plate (AMTI,
185 OR6-7, Watertown, MA, USA). The Ground Reaction Force (GRF) and its point of
186 application were recorded. Kinematic and force plate data were sampled at 60Hz.

187

188 *Data analysis*

189 The following analysis was performed using Matlab (Mathworks, Natick, US).
190 For each trial, the last heel-ground contacts during walking and prior to standing were
191 calculated for left and right foot as the instant of minimum velocity of the toe marker
192 (Pijnappels et al., 2001), and then classified as last heel and penultimate heel contact,
193 irrespective of the side. After the last heel contact, for each trial, the instant of last toe down
194 was identified using the time when the velocity of the toe marker first crossed the zero value.
195 This instant was deemed to be the start of standing, since both feet were on the ground and no
196 further steps were taken (Fig. 1). This instant is called flat feet time, 0s in all mean data
197 figures (Figs 2-4). To accept a trial for analysis, a test was used to confirm that flat feet time
198 represented standing: the vertical component of the ground reaction force had to be within
199 one SD ($\pm 1.7N$) of the value during sustained standing (7.6s later).

200

201 The following variables were calculated.

202 *Upper body velocity and acceleration.* The location of the markers from the pelvis upwards
203 was averaged to provide a representative antero-posterior body location, which was
204 differentiated to provide velocity and acceleration, using a FIR filter (Remez differentiator
205 pass-band filter with pass frequency 1Hz, and stop frequency 6Hz). The whole trial (30s) was
206 then reduced to the correct 7.6s from flat feet time, removing the appropriate initial and final
207 part of the recording and avoiding any filtering distortion at the beginning or end of the trial.

208

209 *Antero-posterior Centre of Gravity (CoG_{AP}) location.* CoG_{AP} was calculated by zero-lag low-
210 pass filtering the sagittal component of the centre of pressure (from point of application) with
211 a cut-off frequency of 0.5Hz (Caron et al., 1997; Loram and Lakie, 2002). This calculation is
212 valid for and presented only during standing. We used this quantity to minimise the possible
213 bias induced by modelling different body shapes and sizes using kinematic data.

214

215 *Antero-posterior CoG position relative to the lower limb joint centres (CoG-Ankle, CoG-
216 Knee and CoG-Hip).* The displacement between a vertical line through the CoG_{AP} and the
217 joint centres was calculated. The joint centre positions were calculated using a combination
218 of surface markers on bony landmarks (Vicon) and MRI imaging (see also above in
219 Apparatus and measurements). The ankle joint centre was calculated relative to the lateral
220 malleolus using the individually measured ankle width. The knee joint centre was calculated
221 as the centre of a line joining markers on lateral and medial femoral epicondyles. The hip
222 joint centre was calculated according to the GaitLab algorithm (Vaughan et al., 1999) using
223 three markers (sacrum, left and right anterior superior iliac spines) and anthropometric
224 measures taken from each subject. For ten participants (three were excluded for
225 contraindication to MRI) the GaitLab calculation was corrected by analysis of the MRI
226 images. The geometrical hip joint centre in the anterior-posterior direction relative to the cod
227 liver oil pills (placed on greater trochanter and iliac spines landmarks) was calculated,
228 approximating the head of the femur as a circular section and assuming its centre as the joint
229 centre (Osirix 2.7.5, Osirix Foundation, Geneva, Switzerland). The joint location used in the
230 kinematic analysis was corrected using each participant's difference between joint calculated
231 from the marker and from the MRI scan. In the sagittal plane, the mean anterior/posterior
232 correction was $\pm 2 \pm 1$ mm (mean \pm SD). For the participants which were excluded from MRI
233 scans, the joint locations were not corrected and the ones calculated using anthropometry and
234 kinematic model were used.

235 Left and right sagittal joint location were averaged. A displacement of 0 mm indicates that
236 the CoG_{AP} is in line with the joint centre.

237

238 For each variable listed above and for the antero-posterior force from the force platform, a
239 time constant was calculated for each trial between flat feet time and 7.6s. The time constant
240 represents the elapsed time for the system response to decay/grow by 1/e at the initial rate.
241 An exponential curve was fitted to the data and the time constant was estimated for each trial
242 and then averaged across participants.

243

244 *Joint moments (MAnkle, MKnee and MHip).* The flexion-extension joint moments were
245 calculated using an inverse dynamic approach (Vaughan et al., 1999). At the ankle, positive
246 values indicate dorsi-flexion moment, while negative values indicate plantar-flexion moment.
247 At the knee and hip, positive values indicate flexion moments.

248

249 For each variable, the mean over three time intervals was calculated. The intervals were
250 chosen to represent the possible phases of standing after walking. The interval duration was
251 determined by a suitable duration that could capture the rapid changes occurring after
252 standing initiation. Therefore, beginning of standing ('Beg') was between flat feet time and
253 0.05s later. This arbitrary choice determined the interval duration, which was kept constant.
254 Steady-state standing ('End') was identified as the latest interval available from the
255 recordings (7.6s), so that interval was between 7.55 and 7.6s. An intermediate interval
256 ('Mid') was selected to start later than the threshold body acceleration time constant (0.25s),
257 but still adequate to record any early changes in configuration. The chosen Mid interval was
258 between 0.5 and 0.55s.

259

260 ***Statistical analysis***

261 A repeated measures univariate ANOVA was run on the CoG_{AP-Joints} (generic term to
262 indicated the displacement between CoG and the leg joints included in this study). Interval (3
263 levels), and Trial (4 levels) were fixed factors and Participant (13 levels) was the random
264 factor. This analysis was conducted using SPSS (ver.24, IBM).

265 Each CoG_{AP-Joint} and joint moments were tested to see if a difference was significant between
266 the two intervals after the deceleration had ceased (Mid vs End) using a two-tailed paired t-
267 test. We used Mid rather than Beg interval in this analysis to measure changes in
268 configuration beyond body deceleration after walking.

269

270 Significance is reported at $p < 0.05$. Unless otherwise stated, results are reported as
271 mean \pm standard error of the mean in the text, and 95% confidence curves are shown in the
272 figures (dotted).

273 **Results**

274 All participants ended their gait with both feet flat on the force plate with a broadly
275 symmetrical, self-chosen stance and foot placement. In Figure 1, a representative participant
276 illustrates the small changes in configuration in a trial and show the need for high precision
277 measurements.

278
279 The transition from walking to standing requires reduction of forward velocity and attainment
280 of equilibrium. Following flat feet time, upper body deceleration and antero-posterior shear
281 force are reduced to the steady-state value rapidly (Fig.2). From all trials, the time constant of
282 the upper body deceleration was 0.16 ± 0.03 s, and a similar time constant was calculated for
283 the shear force 0.15 ± 0.03 s (mean \pm S.D.). These results designated $\tau=0.16$ s as the higher value
284 after which body deceleration had reached a value closer to steady state.

285
286 If the main process governing the joints adjustments is only related to body deceleration, one
287 would expect that the time constant of all the other variables to be close to $\tau=0.16$ s. This is a
288 justified approach considering that body sway in quiet standing determines a not-null mean
289 acceleration (i.e. between -0.031 and $+0.035$ m/s² range measured in the current experiment
290 at steady-state standing). Thus, $\tau=0.16$ s is consistent with the time needed for the body to
291 approximate quiet standing. In order to define a conservative time threshold beyond which
292 the changes measured were not merely related to body deceleration, we used the mean upper
293 body deceleration time constant (0.16) + $3 \times \text{SD}$ (3×0.03), and obtained a value of 0.25 s. Using
294 three times SD gives our analysis 99.7% probability to be investigating adjustments that were
295 not merely linked to body deceleration. If we found higher time constants, we were entitled to
296 investigate the process occurring after 0.25 s.

297

298 **Is body configuration only governed by body deceleration?**

299 Initially, CoG_{AP} was in front of ankle, knee and hip by 38mm, 28mm and 25mm respectively
300 (Fig.3). The hip quickly, $\tau_{\text{hip}}=0.25 \pm 0.12$ s (mean \pm S.D.), aligned more with CoG_{AP} by 27mm
301 (Fig.3B). The displacement between ankle and knee joint centres and CoG_{AP} increased (i.e.
302 misalignment) by 19mm and 13mm progressively with $\tau_{\text{ankle}}=0.62 \pm 0.17$ s and
303 $\tau_{\text{knee}}=0.61 \pm 0.29$ s (mean \pm S.D. Fig.3D,C). These longer time constants suggest a slower
304 process, not related only to the deceleration of the body. The joint moments (Fig.4) showed a
305 similar transition. The ankle and knee moments increased by 6.79Nm and 7.97Nm
306 respectively, while the hip moment decreased by 9.78Nm and transitioned from extension at
307 the beginning of standing to flexion at steady-state standing.

308

309 We investigated the repeatability and trend in CoG_{AP-Joints} and moments.

310 *Univariate analysis.* CoG_{AP-Ankle} showed a significant difference between participants
311 ($F(12,21.857)=2.722$, $p=0.020$), but no difference with trial ($F(3,27)=1.516$, $p=0.233$) or
312 interval ($F(2,24.619)=1.255$, $p=0.303$). An interaction interval \times participant ($F(24,54)=6.962$,
313 $p<0.001$) was found.

314

315 CoG_{AP-Knee} and CoG_{AP-Hip} showed a significant difference between intervals
316 ($F_{\text{Knee}}(2,27.531)=23.707$, $p_{\text{Knee}}<0.001$; $F_{\text{Hip}}(2,33.559)=52490$, $p_{\text{Hip}}<0.001$). CoG_{AP-Knee} and
317 CoG_{AP-Hip} showed no difference between trials ($F_{\text{Knee}}(3,27)=1.557$, $p_{\text{Knee}}=0.223$;
318 $F_{\text{Hip}}(3,27)=0.045$, $p_{\text{Hip}}=0.987$) or participants ($F_{\text{Knee}}(12,14.007)=1.437$, $p_{\text{Knee}}=0.256$;
319 $F_{\text{Hip}}(12,5.713)=0.602$, $p_{\text{Hip}}=0.785$).

320

321 Ankle, knee and hip moment (Fig.4) showed a significant difference between intervals
322 ($F_{\text{Ankle}}(2,30.900)=8.188$, $p_{\text{Ankle}}=0.001$; $F_{\text{Knee}}(2,29.369)=5.601$, $p_{\text{Knee}}=0.009$;

323 $F_{\text{Hip}}(2,27.106)=13.173$, $p_{\text{Hip}}<0.001$). Ankle and knee moment showed a significant difference
324 between participants ($F_{\text{Ankle}}(12,20.887)=7.496$, $p_{\text{Ankle}}<0.001$; $F_{\text{Knee}}(12,13.760)=6.806$,
325 $p_{\text{Knee}}=0.001$; $F_{\text{Hip}}(12,19.353)=0.897$, $p_{\text{Hip}}=0.565$). For none of the joint moments, a difference
326 according to trial was found ($F_{\text{Ankle}}(3,27)=0.089$, $p_{\text{Ankle}}=0.965$; $F_{\text{Knee}}(3,27)=2.567$,
327 $p_{\text{Knee}}=0.075$; $F_{\text{Hip}}(3,27)=1.447$, $p_{\text{Hip}}=0.251$).
328 An interaction trial x participant was also significant for the ankle moment ($F(27,54)=3.875$,
329 $p<0.001$).

330

331 *Two tailed pairwise t-test.* To measure whether the steady-state configuration was resulting
332 only from biomechanical factors, we analysed the intervals after the threshold acceleration
333 time constant (Mid and End). We could not find a difference between body acceleration and
334 velocity between the Mid and End interval (mean \pm sem $\text{vel}_{\text{Mid}}=0.002\pm 0.004\text{m/s}$,
335 $\text{vel}_{\text{End}}=0.003\pm 0.004\text{m/s}$, $p=0.92$; $\text{acc}_{\text{Mid}}=-0.001\pm 0.007\text{m/s}^2$, $\text{acc}_{\text{End}}=0.009\pm 0.005\text{m/s}^2$, $p=0.22$).

336

337 As shown in figure 3E-G, $\text{CoG}_{\text{AP-Ankle}}$ and $\text{CoG}_{\text{AP-Hip}}$ showed a significant difference between
338 the two intervals ($\text{CoG}_{\text{AP-Ankle_Mid}}=47.27\pm 4.08\text{mm}$, $\text{CoG}_{\text{AP-Ankle_End}}=58.33\pm 5.29\text{mm}$,
339 $p_{\text{Ankle}}=0.0062$; $\text{CoG}_{\text{AP-Hip_Mid}}=1.77\pm 5.23\text{mm}$, $\text{CoG}_{\text{AP-Hip_End}}=-5.33\pm 5.21\text{mm}$, $p_{\text{Hip}}=0.0428$).
340 $\text{CoG}_{\text{AP-Knee}}$ did not show a significant difference between the two intervals ($\text{CoG}_{\text{AP-}}$
341 $\text{Knee_Mid}}=25.83\pm 7.61\text{mm}$ $\text{CoG}_{\text{AP-Knee_End}}=32.29\pm 6.89\text{mm}$, $p_{\text{Knee}}=0.0655$).

342

343 As shown in figure 4, the joint moments significantly changed between the two intervals
344 ($p_{\text{Ankle}}=0.0023$; $p_{\text{Knee}}=0.0008$; $p_{\text{Hip}}=0.0001$). The ankle moment changed from
345 $M_{\text{Ankle_Mid}}=17.10\pm 4.36\text{Nm}$ to $M_{\text{Ankle_End}}=26.33\pm 5.26\text{Nm}$. The knee moment changed from
346 $M_{\text{Knee_Mid}}=17.62\pm 5.13\text{Nm}$ to $M_{\text{Knee_End}}=33.98\pm 5.82\text{Nm}$. The hip moment changed from
347 $M_{\text{Hip_Mid}}=12.81\pm 10.50\text{Nm}$ to $M_{\text{Hip_End}}=-31.19\pm 5.54\text{Nm}$.

348 **Discussion**

349 In this study, the tight control of lower limb joint configuration was measured in healthy
350 adults when transitioning to standing after walking. Previous work has demonstrated that
351 neural control is required to maintain the location of the line of gravity with respect to the
352 ankle joint (Loram and Lakie, 2002; Casadio et al., 2005; Loram et al., 2005, 2007; Kiemel et
353 al., 2008, 2011; Morasso and Schieppati, 1999; Mirbagheri et al., 2000). In perturbed
354 standing, also the location of the line of gravity with respect to the hip joint is controlled (see
355 hip strategy, Horak and Nashner 1986). In this study, we measured slow changes in leg joints
356 configuration and repeatable steady-state standing configuration within an individual.
357 Although differences in leg configuration between initiation of standing and steady-state
358 could be expected to decelerate the body after walking, we measured subsequent changes in
359 configuration that are not mechanically required to maintain standing (between Mid and End
360 intervals). Here we discuss the possible reasons for the changes in leg configuration.

361
362 ***Misaligned joint reference in quiet standing***

363 At initiation of standing, we measured fast body deceleration as prompt regulation of
364 acceleration is necessary to remain standing without taking steps after walking. After this
365 deceleration, on average, the leg joints became progressively more misaligned with the
366 vertical (Mid vs End intervals, ankle and hip statistically significant, knee showed a trend).
367 Because we allowed the participants to walk freely in the laboratory before coming to a
368 standing position on the force platform, we can suggest that the observed steady-state
369 misalignment is independent of the body configuration at end of walking. Finally, our
370 analysis could not find a significant difference between the trials performed by the
371 participants. Although a lack of significant difference needs to be cautiously interpreted, the
372 fact that we could not find differences despite intervening experiments and breaks, which are
373 likely to increase variability between trials, may suggest that the data is consistent and that
374 the misalignment is not random.

375
376 It is well known that the ankle is misaligned with the vertical in standing, but the result that
377 also the knee and hip configuration became progressively more misaligned was unexpected
378 and requires further explanations.

379
380 Because misalignment induces a higher external gravitational moment at the joint, there is no
381 mechanical explanation for the transition in configuration observed here. In feedback control
382 theory it is accepted that movement is controlled via a pre-programmed combination of set
383 points, thresholds and feedback gains associated with maintaining or changing a
384 configuration of the body. These ideas are common and have been routinely applied to
385 physiological and postural control (c.f. Bernstein, 1967; Feldman, 1986; Fitzpatrick et al.,
386 1996; Maurer and Peterka, 2005; Lockhart and Ting, 2007; Welch and Ting, 2008). Part of
387 this interpretation is the concept of a set point that the feedback system seeks to maintain or
388 restore following a perturbation. In this experiment, we perturbed human standing by asking
389 participants to walk. We could expect that the body configuration does not change after
390 initiation of standing or, if changes were measured, they were random and not consistent.
391 Instead, we found repeatable changes in configuration, despite different preceding gaits.
392 These changes in configuration suggest that standing was not determined by the end of
393 walking configuration, but other factors were involved in the control of the leg configuration.
394 After gait, the body was in a different configuration. The discrepancy between expected and
395 current position could be monitored and minimised (Bays and Wolpert 2007). We suggest
396 that corrections were made when the relative joint positions were beyond threshold limits, as

397 at initiation of standing. Our hypothesis is that our participants had a body configuration
398 reference which was expected and monitored by the nervous system.

399

400 These results do not preclude the established finding that, during long durations of standing,
401 there would be changes in the reference, for example in response to local irritation, fatigue
402 and need for variation (Duarte and Zatsiorsky, 1999; Duarte and Sternad, 2008). However,
403 within the experiment conditions and although the initial joint configuration was perturbed
404 mainly in the direction of the preceding gait phase, we measured adjustments that drove the
405 leg joints towards the steady-state standing configuration. We can assume that the body
406 configuration measured at steady-state standing is, therefore, an approximation of the body
407 configuration reference in standing. Although this study's conclusions are only congruent
408 with the limited number of trials and short duration of standing analysed, we measured a
409 standing reference which is a misaligned configuration at the lower limb joints.

410

411 *Neural control of leg joints in standing*

412 Investigating how this misaligned standing configuration is maintained is ambitious. Here we
413 can only draw conclusions and propose speculations based on our data.

414

415 We have shown that there is no simple mechanical explanation for the delayed process that
416 we observed between Mid and End intervals. The increased misalignment and joint moments
417 show that steady-state configuration was not necessarily consistent with an energy
418 minimisation/optimisation principle. This poses a key question: Why participants tend to
419 stand in a more misaligned configuration?

420

421 The steady-state misaligned configuration could be consistent with an end of range joint
422 flexion/extension that allows passive stabilisation through joint and ligaments locking (close
423 packed). This configuration allows energy conservation because the congruency between
424 articular surfaces allow load distribution and minimises the energy required to maintain a
425 posture. We could not measure whether the participants maintained a close packed joint
426 position at the end of their flexion/extension range, but our results show that the misaligned
427 configuration could be achieved through modulation in joint moments (Fig. 4) which allows
428 small body sway around an average position. This possibility is consistent with
429 proprioceptive mechanisms of standing.

430

431 In standing, joint positions have to be sensed, otherwise internal and external perturbations
432 may destabilise the body and lead to loss of balance. It is unclear whether proprioception of
433 small, postural joint rotations is improved by lower modulation of muscular activity (Hulliger
434 et al., 1982; Cody et al., 1986; Di Giulio et al., 2009; Loram et al., 2009) or by slight tonic
435 activity (Fitzpatrick and McCloskey, 1994). However, proprioception is ambiguous when
436 sensing absolute position, rather than its change (Proske and Gandevia, 2012). On the other
437 hand, the nervous system is exceptionally sensitive to central estimation of muscle forces and
438 movement responses to maintain equilibrium (Fitzpatrick and McCloskey, 1994). In this
439 framework, muscle activation involved in modulating joint moments provides an estimate of
440 the mean body configuration. The configuration thresholds and reference could be coded in
441 terms of muscle activation patterns. Our hypothesis is that the muscle activation at a certain
442 point in time could be compared to the reference activation patterns, and muscle activation
443 would be modulated to facilitate standing control. We suggest that this mechanism may be
444 involved in quiet standing.

445

446 It is noteworthy that differences could be seen between participants. Investigating these
447 differences and their functional implications is beyond the scope of this study. However, it is
448 possible that particular training techniques, injuries or compensatory mechanisms may be at
449 the basis of this kind of differences, and the reference muscular activation pattern could be
450 different between individuals.

451

452 **Limitations**

453 In this study, we did not find a statistical difference between trials and we suggest that this
454 may confirm that the misalignment measured is consistent within a participant. However, the
455 number of trials available per participant varied between 2 and 4 due to technical problems
456 that were only discovered during post-processing. We have interpreted this result cautiously,
457 but the fact that no differences were found despite an intervening long break led us to
458 conclude that consistency and repeatability of the data is acceptable. Furthermore, we chose
459 to treat Trial as a fixed factor. One may consider Trial as a random factor because there is no
460 meaningful, consistent difference between the levels. However, in order to consider Trial as a
461 random factor, Trial needs to be an instance from a large number of repetitions that have been
462 conducted, chosen at random from a larger subset of similar repetitions. This was not the case
463 in the current study because there was an intervening experiment and this is the reason of our
464 statistical model set-up.

465

466 Based on the measurements and analyses conducted, we suggest that other factors rather than
467 passive and biomechanical factors are involved in standing. It is difficult to distinguish
468 between active and passive mechanisms at the transition between initiation and steady-state
469 standing, particularly because of the body inertia and the possible non-linear muscle
470 behaviour during the transition. This experiment was designed to test whether biomechanical
471 and passive mechanisms could fully explain body position transition between initiation and
472 steady-state standing. In the analysis used here, we aimed to measure changes beyond body
473 deceleration and inertia. This is the reason why we calculated the time constant of body
474 acceleration and we used $3 \times SD$ and we reported changes between the 'Mid' and the 'End'
475 intervals, rather than the 'Beg' interval. Despite this analysis, one limitation is that other non-
476 active mechanisms may be still involved in the transition, but here we suggest that the
477 increased misalignment and joint moments at steady-state standing cannot exclude an active
478 control of configuration. Further experiments are needed to confirm this suggestion, but in
479 this study we were able to use an ecological protocol and measure physiological mechanisms
480 that are consistent with the hypothesis of active knee and hip control in quiet standing.

481

482

483 **Conclusions**

484 In this study, we measured leg joint configuration in standing after walking as a way to
485 physiologically perturb this configuration. We found that the misalignment between line of
486 gravity and ankle, knee and hip and the joint moments were larger at steady-state standing.
487 We, therefore, suggest that the human leg joints are controlled relative to a misaligned
488 standing reference. Although the experimental data presents limitations due to trial duration
489 and number of trials, we measured increased joint moments between two intervals after
490 initiation of standing (Mid and End). Because there is no need for a modulation of knee and
491 hip moments in quiet standing, our results suggest that muscle moments were modulated to
492 achieve and maintain the steady-state standing configuration. Additional work is needed to
493 support the current evidence, possibility involving modelling of this tight control in standing.
494 Here we suggest that modulation of joints moments constitutes an additional voluntary

495 control mechanism, other than the well-established passive and tonic control mechanisms,
496 involved in maintaining quiet standing in humans.
497

498 **Figure Legends**

499 **Figure 1. Representative trial.**

500 From top to bottom, whole body sagittal stick figure from markers location of representative
501 participant at 0.5s before flat feet time, flat feet time and 2.5s, 5s and 10s after flat feet time.
502 Vertical dashed line represents line of gravity location.
503 For the representative trial, relative displacement between CoG and hip, knee and ankle
504 position between 0.5s before flat feet time and end of trial. Vertical dashed line represents flat
505 feet time, dotted line identify the intervals for which differences were calculated in the
506 analysis.

507

508 **Figure 2. Whole body quantities.**

509 From top to bottom, mean (solid) \pm 95% confidence intervals (dotted) of upper body antero-
510 posterior velocity (A), deceleration (B) and shear force from the force platform (C). Body
511 velocity was quickly reduced to approximate the end of standing velocity.
512 Vertical dashed lines illustrate the three intervals used in the statistical analysis: beginning of
513 standing ('Beg' from flat feet time $t=0s$ to 0.05s later), after body deceleration have ceased
514 ('Mid' 0.5-0.55s after flat feet time) and steady-state standing ('End', 7.55-7.6s after flat feet
515 time).

516

517 **Figure 3. Antero-posterior leg joint position relative to centre of gravity position.**

518 From top to bottom, (A) mean stick figure obtained from sagittal position of lower body
519 markers' locations (toe and sacrum) and calculated joint centres (ankle, knee and hip) relative
520 to line of gravity location (dashed) from all the trials at five time points (flat feet time, 0.5s,
521 2.5s, 5s and 7.6s after flat feet time). Mean (solid) \pm 95% confidence intervals (dotted) of hip
522 (B), knee (C) and ankle (D) joint centre location relative to centre of gravity position for the
523 common duration to all the trials included in the analysis (i.e. 7.6s). 0m represents perfect
524 sagittal alignment between joint centre and gravity.

525 Vertical dashed lines illustrate the three intervals used in the statistical analysis: beginning of
526 standing ('Beg' from flat feet time $t=0s$ to 0.05s later), after body deceleration have ceased
527 ('Mid' 0.5-0.55s after flat feet time) and steady-state standing ('End', 7.55-7.6s after flat feet
528 time).

529 Mean \pm standard error of the mean at Mid and End intervals for CoG-Hip (E), CoG-Knee (F),
530 CoG-Ankle (G). One star indicates $p<0.05$, two stars $p<0.01$.

531

532 **Figure 4. Sagittal leg joint moments.**

533 From top to bottom, mean (solid) \pm 95% confidence intervals (dotted) of hip (A), knee (B)
534 and ankle (C) sagittal moments for the common duration to all the trials included in the
535 analysis (i.e. 7.6s).

536 Vertical dashed lines illustrate the three intervals used in the statistical analysis: beginning of
537 standing ('Beg' from flat feet time $t=0s$ to 0.05s later), after body deceleration have ceased
538 ('Mid' 0.5-0.55s after flat feet time) and steady-state standing ('End', 7.55-7.6s after flat feet
539 time).

540 Mean \pm standard error of the mean at Mid and End interval for hip flexion-extension moment
541 (D), knee flexion-extension moment (E), and ankle dorsi-plantarflexion moment (F). Two
542 stars indicate $p<0.01$, three stars $p<0.001$.

543

544

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549

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551 All authors contributed to design of the work and critically reviewed the intellectual content.
552 IDG contributed to acquisition, analysis and drafting of the work. IDG and VB contributed to
553 manuscript writing. The authors approved the final version of the manuscript and agree to be
554 accountable for all aspects of the work in ensuring that questions related to the accuracy or
555 integrity of any part of the work are appropriately investigated and resolved.

556

557 **Conflict of Interest Statement**

558 The submitted work was carried out in absence of any personal, professional or financial
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560

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563

564 **Data Availability Statements**

565 The raw data supporting the conclusions of this manuscript will be made available by the
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567

568

569 **References**

- 570 Bays PM, Wolpert DM & Flanagan JR (2005). Perception of the consequences of self-action
571 is temporally tuned and event driven. *Curr Biol* 15(12):1125–1128.
572
- 573 Bernstein N. (1967). *The co-ordination and regulation of movements*. Pergamon Press Ltd.,
574 London.
575
- 576 Caron O, Faure B, Breniere Y (1997). Estimating the centre of gravity of the body on the
577 basis of the centre of pressure in standing posture. *J Biomech* 30(11/12):1169-1171.
578
- 579 Casadio M, Morasso PG & Sanguineti V (2005). Direct measurement of ankle stiffness
580 during quiet standing: implications for control modelling and clinical application. *Gait*
581 *Posture* 21(4):410-424.
582
- 583 Cody FWJ, Macdermott N, Matthews PBC & Richardson HC. (1986). Observations on the
584 genesis of the stretch reflex in Parkinson's disease. *Brain* 109: 229-249.
585
- 586 Duarte M & Zatsiorsky VM (1999). Patterns of center of pressure migration during prolonged
587 unconstrained standing. *Motor Control* 3(1):12-27.
588
- 589 Duarte M & and Sternad D (2008). Complexity of human postural control in young and older
590 adults during prolonged standing. *Exp Brain Res* 191(3):265-276.
591
- 592 Di Giulio I, Baltzopoulos V, Managanaris CN & Loram ID (2013). Human standing: does the
593 control strategy pre-program a rigid knee? *J Appl Physiol* 114(12):1717-1729. DOI:
594 10.1152/jappphysiol.01299.2012.
595
- 596 Di Giulio I, Maganaris CN, Baltzopoulos V & Loram ID. (2009). The proprioceptive and
597 agonist roles of gastrocnemius, soleus and tibialis anterior muscles in maintaining human
598 upright posture. *J Physiol* 587:2399-2416.
599
- 600 Feldman AG & Levin MF. (2009). The equilibrium-point hypothesis - Past, present and
601 future. In *Progress in motor control: A multidisciplinary perspective*, ed. Sternad D &
602 University PS, pp. 699-726. Springer US, New York.
603
- 604 Fitzpatrick R & McCloskey DI (1994). Proprioceptive, visual and vestibular thresholds for
605 the perception of sway during standing in humans. *J Physiol* 478 (1):173-186.
606
- 607 Fitzpatrick R, Burke D & Gandevia SC (1996). Loop gain of reflexes controlling human
608 standing measured with the use of postural and vestibular disturbances. *J Neurophysiol*
609 76(6):3994-4008.
610
- 611 Gatev P, Thomas S, Kepple T & Hallett M (1999). Feedforward ankle strategy of balance
612 during quiet stance in adults. *J Physiol* 514(3):915-928.
613
- 614 Horak FB, Nashner LM (1986). Central programming of postural movements: adaptation to
615 altered support-surface configurations. *J Neurophysiol* 55(6): 1369-1381.
616

617 Hulliger M, Nordh E & Vallbo AB. (1982). The absence of position response in spindle
618 afferent units from human finger muscles during accurate position holding. *J Physiol*
619 322:167-179.
620

621 Hsu W-L, Scholz JP, Schoener G, Jeka JJ, Kiemel T (2007). Control and Estimation of
622 posture during quiet stance depends on multijoint coordination. *J Neurophysiol* 97:3024-
623 3035.
624

625 Kiemel T, Elahi AJ & Jeka JJ (2008). Identification of the plant for upright stance in humans:
626 multiple movement patterns from a single neural strategy. *J Neurophysiol* 100(6):3394–3406.
627 DOI: 10.1152/jn.01272.2007.
628

629 Kiemel T, Zhang Y & Jeka JJ (2011). Identification of Neural Feedback for Upright Stance in
630 Humans: Stabilization Rather Than Sway Minimization. *J Neurosci* 31(42):15144–15153.
631 DOI: 10.1523/JNEUROSCI.1013-11.2011.
632

633 Loram ID & Lakie M (2002). Direct measurement of human ankle stiffness during quiet
634 standing: the intrinsic mechanical stiffness is insufficient for stability. *J Physiol* 545(3):1041–
635 1053.
636

637 Loram ID, Lakie M, Di Giulio I & Maganaris CN. (2009). The consequences of short-range
638 stiffness and fluctuating muscle activity for proprioception of postural joint rotations: the
639 relevance to human standing. *J Neurophysiol* 102: 460-474.
640

641 Loram ID, Maganaris CN & Lakie M (2005). Active, non-spring-like muscle movements in
642 human postural sway: how might paradoxical changes in muscle length be produced? *J*
643 *Physiol* 564(1):281–293.
644

645 Loram ID, Maganaris CN & Lakie M (2007). The passive, human calf muscles in relation to
646 standing: the non-linear decrease from short range to long range stiffness. *J Physiol*
647 584(2):661–675.
648

649 Lockhart DB & Ting LH. (2007). Optimal sensorimotor transformations for balance. *Nat*
650 *Neurosci* 10:1329-1336.
651

652 MacConaill MA & Basmajian JV. (1977). *Muscles and movements. A basis for human*
653 *kinesiology*. Robert E. Krieger Publishing Co, Inc., New York.
654

655 Maurer C & Peterka RJ. (2005). A new interpretation of spontaneous sway measures based
656 on a simple model of human postural control. *J Neurophysiol* 93:189-200.
657

658 Mirbagheri MM, Barbeau H & Kearney RE (2000). Intrinsic and reflex contributions to
659 human ankle stiffness: variation with activation level and position. *Exp Brain Res*
660 135(4):423-436.
661

662 Morasso PG & Schieppati M (1999). Can muscle stiffness alone stabilize upright standing? *J*
663 *Neurophysiol* 82(3):1622-1626.
664

665 Peterka RJ & Benolken MS (1995). Role of somatosensory and vestibular cues in attenuating
666 visually induced human postural sway. *Exp Brain Res* 105(1):101-110.

667
668 Pijnappels M, Bobbert MF & van Dieen JH. (2001). Changes in walking pattern caused by
669 the possibility of a tripping reaction. *Gait Posture* 14(1):11-18.
670
671 Proske U & Gandevia SC (2012). The proprioceptive senses: their roles in signaling body
672 shape, body position and movement, and muscle force. *Physiol Rev* 92(4):1651–1697.
673
674 Steindler A. (1964). Kinesiology of the human body under normal and pathological
675 conditions. Charles C. Thomas Publisher, Springfield, Illinois, USA.
676
677 Vaughan CL, Davis BL & O'Connor JC. (1999). *Dynamics of human gait*. Kiboho
678 Publishers, Western Cape.
679
680 Welch TDJ & Ting LH. (2008). A feedback model reproduces muscle activity during human
681 postural responses to support-surface translations. *J Neurophysiol* 99:1032-1038.
682