Asymmetric gait patterns alter the reactive control of intersegmental coordination patterns in the sagittal plane during walking

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Abstract

Recovery from perturbations during walking is primarily mediated by reactive control strategies that coordinate multiple body segments to maintain balance. Balance control is often impaired in clinical populations who walk with spatiotemporally asymmetric gait, and, as a result, rehabilitation efforts often seek to reduce asymmetries in these populations. Previous work has demonstrated that the presence of spatiotemporal asymmetries during walking does not impair the control of whole-body dynamics during perturbation recovery. However, it remains to be seen how the neuromotor system adjusts intersegmental coordination patterns to maintain invariant whole-body dynamics. Here, we determined if the neuromotor system generates stereotypical coordination patterns irrespective of the level of asymmetry or if the neuromotor system allows for variance in intersegmental coordination patterns to stabilize whole-body dynamics in the sagittal plane. Nineteen healthy participants walked on a dual-belt treadmill at a range of step length asymmetries, and they responded to unpredictable, slip-like perturbations. We used principal component analysis of segmental angular momenta to characterize intersegmental coordination patterns before, during, and after imposed perturbations. We found that two principal components were sufficient to explain \(~95\%\) of the variance in segmental angular momentum during both steady-state walking and responses to perturbations. Our results also revealed that walking with asymmetric step lengths led to changes in intersegmental coordination patterns during the perturbation and during subsequent recovery steps without affecting whole-body angular momentum. These results suggest that the nervous system allows for variance in segment-level coordination patterns to maintain invariant control of whole-body angular momentum during walking. Future studies exploring how these segmental coordination patterns change in individuals with asymmetries that result from neuromotor impairments can provide further insight into how the healthy and impaired nervous system regulates dynamic balance during walking.
1. Introduction

Bipedal locomotion is inherently unstable due to the small base of support, long single-limb support times, and sensorimotor transmission delays [1]. As a result, we must frequently generate corrective responses to maintain balance in response to both internal and external perturbations [2,3]. For example, to recover from unexpected perturbations such as slips or trips while walking, the nervous system generates reactive control strategies involving simultaneous, coordinated responses of both the upper and lower limbs [4,5]. These reactive, interlimb responses to perturbations can restore stability by generating changes in angular momentum that counteract the body’s rotation toward the ground.

One conventional method to capture whole-body rotational dynamics during perturbation responses is to compute whole-body angular momentum (WBAM). WBAM reflects the net influence of all the body segments’ rotation and translation relative to a specified axis, which is commonly taken to project through the body’s center of mass [6–8]. WBAM is highly regulated as its value remains close to zero during normal, unperturbed walking [9,10]. During perturbed walking, angular momentum dramatically deviates from that measured during unperturbed walking [6,7], and this deviation captures the features of body rotation that, if not arrested, would lead to a fall. To regain balance when encountering unexpected perturbations, the central nervous system activates muscles to accelerate body segments and restore angular momentum across multiple recovery steps [11,12].

Angular momentum can also capture balance impairments in populations with gait asymmetries and sensorimotor deficits such as amputees and stroke survivors. These individuals often have a higher peak-to-peak range of angular momentum than healthy controls [13–16], and the presence of gait asymmetries may contribute to balance impairments in these populations. An important question for clinical researchers is whether there is a causal relationship between gait asymmetry and the ability to maintain balance in response to perturbations during walking. Previous work demonstrated that whole-body dynamics, as measured by WBAM, do not change in response to imposed gait asymmetries in healthy individuals [7]. However, the strategy that the central nervous system uses to stabilize whole-body dynamics remains to be determined.

There are two distinct hypotheses capable of explaining the negligible influence of asymmetry on whole-body angular momentum. First, the central nervous system may generate stereotypical, invariant intersegmental coordination patterns in response to perturbations, irrespective of the level of asymmetry. Alternatively, the nervous system could use reactive control strategies that covary with asymmetry in a manner that would lead to invariant control of whole-body momentum. This would be consistent with the uncontrolled manifold (UCM) hypothesis, which proposes that the nervous system allows for variability in segmental angular momenta to stabilize a higher-order performance variable such as whole-body angular momentum [17].

Dimensionality reduction techniques, such as principal component analysis (PCA), are commonly used to capture how the central nervous system coordinates multiple limb segments [6,9,18]. PCA reduces the high-dimensional, multi-segmental time series data into a lower-dimensional set of latent variables capable of capturing the overall variance in angular momentum. For example, two recent studies [6,19] used PCA to quantify intersegmental coordination patterns during unperturbed walking and in response to a set of multi-directional perturbations. The authors found that the over 90% of the variance in segmental angular momentum among 15 body segments could be explained with three to four principal components. Moreover, they found that the coordination patterns observed during perturbation recovery were highly correlated with the patterns observed during unperturbed walking. These
results suggest that PCA can be used to characterize intersegmental coordination during both unperturbed and perturbed gait.

Here, our objective was to determine how the presence of step length asymmetries influences patterns of intersegmental coordination during slip-like perturbations. Since it has previously been demonstrated that step length asymmetry does not influence the magnitude of whole-body angular momentum, we aimed to determine if this was because the neuromotor system generates stereotypical intersegmental coordination patterns across levels of asymmetry or because the neuromotor system generates patterns of intersegmental coordination that covary with spatiotemporal asymmetry. Ultimately, our findings extend our understanding of how the healthy central nervous system coordinates intersegmental dynamics to maintain balance during walking.

2. Methods

2.1. Participant characteristics

A total of 19 healthy young individuals (10M, 24 ± 4 years old) with no musculoskeletal or gait impairments participated in this study. Lower limb dominance was determined by asking participants which leg they would use to kick a ball. The study was approved by the Institutional Review Board at the University of Southern California, and all participants provided informed consent before participating. All aspects of the study conformed to the principles described in the Declaration of Helsinki.

2.2. Experimental protocol

Data used here were collected as part of a previous study [7], and we provide a summary of the procedures and setup below. Participants walked on an instrumented, dual-belt treadmill with force plates underneath (Bertec, USA) at 1.0 m/s for six separate trials and reacted to accelerations of the treadmill belts throughout the experiment. Although 1 m/s was slower than the reported average self-selected speed during treadmill walking [20], we chose this speed to be consistent with other investigations of the role of asymmetry during healthy gait [21–23]. For the first trial, participants walked on the treadmill for three minutes (Baseline) to obtain their natural level of step length asymmetry. Then, for subsequent trials, participants were instructed to modify their step lengths according to visual feedback provided via a display attached to the treadmill, and we informed them that random slip-like perturbations would occur during these trials. The visual feedback displayed the target step length for both right and left legs. A “success” message would appear on the screen if the participants were able to step within three standard deviations of the target step length. The standard deviation was determined on an individual basis from each participant’s baseline step length variability. Participants completed a randomized sequence of five, six-minute trials with target step length asymmetries (SLA, Eq 1) of 0%, ±10%, and ±15% where 0% represents each participant’s baseline SLA.

\[
SLA = \frac{100 \times (SL_{left} - SL_{right})}{SL_{left} + SL_{right}}
\]

SL_{left} represents left step length and SL_{right} represents the right step length. Each trial consisted of one-minute of practice walking without any perturbations, and then a total of 20 perturbations were applied (10 to each belt) during the remainder of the trial. Foot strike was computed as the point when vertical ground reaction forces reached 150 N. Each perturbation was remotely triggered by Python code and was characterized by a trapezoidal speed profile in
which the treadmill accelerated at foot strike to 1.5 m/s at an acceleration of 1.6 m/s², held this
speed for 0.3 s, and then decelerated back to 1.0 m/s at a deceleration of 1.6 m/s² during the
swing phase of the perturbed leg. Participants were aware that they would experience perturba-
tions during the experiment, but the perturbations were randomly triggered to occur within a
range of 20 to 30 steps after the previous perturbation to prevent participants from precisely
anticipating perturbation timing. This range of steps was also selected to provide participants
with sufficient time to reestablish their walking pattern to match with the visual feedback.

2.3. Data acquisition
A ten-camera motion capture system (Qualisys AB, Gothenburg, Sweden) recorded 3D
marker kinematics at 100 Hz and ground reaction forces at 1000 Hz. We placed a set of 19 mm
spherical markers on anatomical landmarks to create a 13-segment, full-body model [24,25].
We placed marker clusters on the upper arms, forearms, thighs, shanks, and the back of heels.
Marker positions were calibrated during a five-second standing trial at the beginning of each
trial. We removed all joint markers after the calibration.

2.4. Data processing
We post-processed the kinematic and kinetic data in Visual3D (C-Motion, Rockville, MD,
USA) and Matlab 2017a (Mathworks, USA) to compute variables of interest. Marker positions
and ground reaction forces were low-pass filtered by 4th order Butterworth filters with cutoff
frequencies of 6 Hz and 20 Hz, respectively. We selected the type of filter and cut-off frequency
based on previous literature [3,26,27]. We calculated the achieved SLA as follows: first, we cal-
culated the mean SLA of the four strides before each perturbation and then distributed these
mean values into five equally spaced bins centered at -15%, -10%, 0, 10%, 15% with bin width
equal to 5%. We used this achieved SLA instead of target SLA as the independent variable in
our statistical analyses. We categorized Baseline (BSL) steps as the two steps before the pertur-
bation occurred, perturbation (PTB) steps as the step during which the perturbation was
applied, and recovery (REC) steps as the four steps that followed the perturbation. Since we
did not find any differences between left and right perturbations, our current analysis includes
only perturbations of the right limb [7]. We also focused our analysis on angular momentum
in the sagittal plane as this was the direction in which the most prominent changes in WBAM
were observed. Only minor deviations in WBAM in the frontal and transverse plane occurred
during the perturbation and recovery steps [7].

2.5. Segmental angular momentum and whole-body angular momentum
We created a 13-segment, whole-body model in Visual3D and calculated the angular momen-
tum of each segment about the body’s center of mass. Segmental angular momenta (Lᵢ) cap-
tured how the rotational and translational behavior of each body segment changed in response
to the treadmill perturbations. The model included the following segments: head, thorax, pel-
vis, upper arms, forearms, thighs, shanks, and feet. The limb segments’ mass was modeled
based on anthropometric tables [28], and segment geometry was modeled based on the
description in Hanavan [29]. All segments were modeled with six degrees of freedom, and we
did not define any constraints between segments. Segmental linear and angular velocity were
computed using Eq 2 [15].

\[ L_i = \frac{m_i (\vec{r}_{CM,i} \times \vec{v}_{CM,i}) + I\vec{\omega}}{MVH} \] (2)

Here, \( m_i \) is segmental mass, \( \vec{r}_{CM,i} \) is a vector from the segment’s COM to the body’s COM,
Where $v_{CM,i}$ is the velocity of each segment’s COM relative to the body’s COM, $I^i$ is the segmental moment of inertia, $\omega^i$ is segmental angular velocity, and the index $i$ corresponds to individual limb segments.

Whole-body angular momentum was calculated as the sum of all segmental angular momenta using Eq 3. In addition, integrated whole-body angular momentum was computed as the area under the curve of the WBAM trajectory for each step cycle to quantify the degree to which the body rotates about its center of mass across a step cycle.

$$\vec{L} = \sum_i L^i_i$$

Lastly, we normalized momentum by the participant’s mass ($M$), baseline treadmill velocity ($V$), and the participant’s height ($H$) following previous literature [9,16]. Since our statistical analysis used a within-subject design, the choice of variables used for normalization should not affect the statistical results. The convention for measuring angular momentum was defined such that positive values represented backward rotation.

### 2.6. Principal component analysis (PCA)

We used principal component analysis (PCA) to extract intersegmental coordination patterns for each step cycle. Before performing PCA, we first time normalized the time series of segmental angular momenta to 100 points for each step cycle. Then, for each participant, we generated an $L_s$ matrix for each achieved SLA ($\pm 15\%$, $\pm 10\%$, $\pm 5\%$, $\%0$) and step type (BSL1, BSL2, PTB, REC1, REC2, REC3, REC4) with $n_{steps}$’100 rows and 13 columns. On average, we created 6 (achieved SLA) by 7 (step types) matrices per participant as not all participants achieved each desired level of asymmetry. We then standardized each matrix to have zero mean and performed PCA to extract subject-specific coordination patterns using the `pca` function in Matlab’s Statistical and Machine Learning Toolbox. Using PCA, we decomposed the segmental angular momenta data into 1) a weighting coefficient matrix consisting of principal components (PCs) ordered according to their variance accounted for (VAF) and 2) time series scores which represented the activation of each PC throughout the step cycle (Fig 1). We retained the number of PCs necessary to account for at least 90% of variance in $L_s$.

### 2.7. Comparison of intersegmental coordination patterns

To investigate how intersegmental coordination patterns changed after each perturbation, we compared the PCs extracted from the perturbation and recovery steps to the PCs extracted from baseline steps. We computed the included angle ($\theta_{step}$, Eq 4) between each pair of PCs as this is a common method to compare the similarity between vectors in a high-dimensional space. The included angle of the unit vectors was between 0˚ (parallel and identical) and 90˚ (orthogonal and most dissimilar) [30].

$$\theta_{step} = \cos^{-1}(\langle PC_{baseline} \cdot PC_{post} \rangle)$$

We then determined if the included angle between perturbation steps and baseline steps was outside the distribution of included angles observed during unperturbed baseline walking. To this end, we performed a permutation test that randomly and repeatedly selected two groups of ten baseline steps for each participant. For each permutation, we first performed PCA for each group of 10 steps and then calculated the included angle between the two PCs. We repeated this shuffling process 10000 times for each participant. We used the median of this distribution as the reference level in the statistical model to determine if the included angle for post-perturbation values was greater than what would be expected from step-to-step variance.
Similarly, we computed the included angle between PCs extracted during walking at different levels of asymmetry to those extracted from symmetrical walking to investigate how
asymmetry influenced intersegmental coordination patterns. (Eq 5).

\[ \theta_{\text{asy}} = \cos^{-1}(\overrightarrow{PC_{\text{sym}}} \cdot \overrightarrow{PC_{\text{asy}}}) \]  

(5)

We also determined if the differences in coordination observed during walking with different levels of asymmetry were above the level of variance observed during symmetrical walking. As described above, we obtained a reference distribution of included angles from symmetric walking to determine if the included angle for each level of asymmetry was greater than what would be expected from natural, step-to-step variance.

2.8. Statistical analysis

All statistical analyses were performed in R (3.4.3) using linear mixed-effects (LME) models. We used the lme4 package to fit the model, the multcomp comparison for multiple comparisons [31], and lmerTest package to calculate p-values [32]. Residual normality was confirmed using the Shapiro-Wilk test. When computing p-values, we used the Satterthwaite approximation for the degrees of freedom based on differences in variance between conditions. We used the Bonferroni correction for multiple comparisons for all post-hoc analyses. For each model, we determined if a model with random effects provided a better fit than a model with only fixed effects by comparing both models using a likelihood ratio test and choosing the most parsimonious model. Based on this analysis, we ultimately included random effects in all models to account for the individual differences between subjects. Significance was set at \( p < 0.05 \) level.

We first determined if the PCs extracted from the recovery steps differed from the PCs extracted from the baseline steps during symmetrical walking. Here, the independent variable was step type, and the dependent variable was \( \theta_{\text{step}} \). The models were fit for both PC1 and PC2. We performed a log transformation of the dependent variable \( \theta_{\text{step}} \) to ensure that the residuals were normally distributed. The full model structure was defined as follows: \( \log(\theta_{\text{step}}) \sim 1 + \text{Step}_\text{Type} + (1|\text{Subject}) \). To provide further insight into how the weighting coefficients for individuals segments varied across steps, we fit linear mixed effect models for each segment to determine if the segment’s weighting coefficients \( (W_{\text{segment}}) \) during perturbation and recovery steps differed from the baseline steps. For this analysis, the independent variable was step type and the dependent variable was each weighting coefficient. The model was fit for each segment and for both PC1 and PC2. The reference level was set to be the right baseline step (BSL1) for all right steps and the left baseline step (BSL2) for all left steps. The full model structure was defined as follows: \( W_{\text{segment}} \sim 1 + \text{Step}_\text{Type} + (1|\text{Subject}) \).

Then, we determined if intersegmental coordination patterns during asymmetrical walking differed from those during symmetrical walking. For this analysis, we used Welch’s t-test to evaluate if the included angle between the PCs extracted from the asymmetrical trials and those extracted from symmetric walking were greater than what would be expected by chance. We used Welch’s t-test because the included angle was not normally distributed.

Lastly, we determined if the included angle between each asymmetric trial and symmetric walking varied with the magnitude or direction of asymmetry. For this analysis, the independent variables were the magnitude of asymmetry \( (|\text{Asym}|) \), the direction of asymmetry \( (\text{Direction}) \), and the interaction between asymmetry magnitude and direction, and the dependent variable was \( \theta_{\text{asy}} \). We performed a log transformation of the dependent variable \( \theta_{\text{asy}} \) to ensure that the residuals were normally distributed. The full model structure was defined as follows: \( \log(\theta_{\text{asy}}) \sim 1 + |\text{Asym}| + \text{Direction} + (1|\text{Subject}) \). We fit separate linear mixed-effect models for each of five steps (Baseline1, Baseline2, Perturbation, Recovery 1 and Recovery 2) and each PC.
3. Results

For all steps, two principal components accounted for ~95% of the variance in segmental angular momentum (Table 1). On average, PC1 explained 74 ± 4% of the variance, and PC2 explained 22 ± 1% of the variance, while PC3 accounted for less than 3% of the variance. Thus, the remaining analysis focuses on the first two PCs.

3.1. Patterns of intersegmental coordination when walking with equal step lengths

Contributions from the lower extremities were typically dominant in the first PC, while contributions from the arms, pelvis, thorax, and head were less prominent (Fig 2). During right steps, the left leg was in the swing phase and generated more positive momentum about the body’s COM, while the right leg generated negative momentum. Thus, the weighting coefficients for the left leg segments (left thigh, shank, and foot) were positive while the coefficients for the right leg segments were negative. Similarly, during a left step, the right leg was in the swing phase and generated more positive momentum about COM, while the left leg generated negative momentum. Thus, the weighting coefficients were positive while the coefficients for the left leg segments were negative. Overall, the first PC captured the opposing momenta of the two legs resulting from differences in the direction of rotation relative to the body's center of mass.

For PC2, weighting coefficients for distal segments were also larger than the weighting coefficients for proximal segments, although the coefficient for the thorax (THX) increased compared to that in PC1. During the right step, the left thigh and left shank’s momenta opposed the momentum of the left foot. Similarly, during the left step, the right thigh and shank momenta opposed the right foot momentum. Thus, PC2 captured intralimb cancellation of segmental momenta.

3.2. Effects of perturbations on patterns of intersegmental coordination

During the perturbation step, there was a significant increase in the included angle compared with the reference estimated of step-to-step variance, which indicated that the intersegmental coordination patterns during perturbation steps differed from the coordination patterns during baseline steps (Fig 3). For PC1, we found that the intersegmental coordination patterns were significantly different from the patterns during baseline walking for the perturbation steps (t(80) = 18.2, p<2e-16), first recovery steps (t(80) = 11.8, p<2e-16), and second recovery steps (t(80) = 8.4, p = 2.2e-8). Similarly, for PC2, intersegmental coordination differed during perturbation steps (t(80) = 10.8, p<2.0e-16), first recovery steps (t(80) = 12.5, p<2e-16), second recovery steps (t(80) = 4.3, p = 3.81e-5), and fourth recovery step (t(80) = 2.3, p = 0.02). There was no significant difference between intersegmental coordination patterns during the third recovery steps for either PC1 (p = 0.58) or PC2 (p = 0.22). Thus, participants generally were able to restore their coordination patterns to baseline by the third recovery step.

Table 1. Variance accounted for (VAF) for PC1, PC2, and PC3 during baseline steps, perturbation steps, and recovery steps.

| Step Type       | PC1     | PC2     | PC3     | Sum     |
|-----------------|---------|---------|---------|---------|
| Baseline steps  | 74±4%   | 22±5%   | 2±1%    | 98±1%   |
| Perturbation steps | 75±5%   | 20±5%   | 3±1%    | 98±1%   |
| Recovery steps  | 74±3%   | 21±4%   | 3±1%    | 98±1%   |
| All steps       | 74±4%   | 22±4%   | 2±1%    | 98±1%   |

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We also observed a significant main effect of step type on segment weighting coefficients for PC1 for all segments (all \( p < 0.005 \)) except the pelvis (\( p = 0.02 \)). Similarly, for PC2, we found a significant main effect of step type on segment weighting coefficients for all segments (all \( p < 0.005 \)).

### 3.3. Effects of step length asymmetry on patterns of intersegmental coordination

Although the general patterns of intersegmental coordination were similar across levels of asymmetry, asymmetric walking patterns led to measurable changes in the contributions of the distal lower extremity segments (Fig 4).

As the magnitude of achieved asymmetry increased, we observed an increase in the deviation of intersegmental coordination patterns from symmetrical walking (Fig 5A and 5B) while there was no change in integrated whole-body angular momentum compared to symmetrical walking (Fig 5C). One outlier was removed before fitting the linear mixed model for the perturbation step for PC2 because it was more than three standard deviations higher than the median of the included angles. Excluding the outlier did not change the statistical outcome. All included angles differed from the permuted estimate of included angles (\( p < 0.05 \)), indicating that intersegmental coordination at each level of asymmetry differed from the coordination pattern during symmetrical walking. For all steps, we observed a significant main effect of asymmetry on the included angle between the PCs from the asymmetric trials and the symmetric trial (Table 2).

The included angle between the PCs extracted during asymmetric walking and symmetric walking increased with the magnitude of achieved asymmetry (Fig 5A and 5B). Specifically,
the difference between intersegmental coordination patterns was greater when walking with 15% asymmetry compared to 5% asymmetry during right baseline steps (Bonferroni corrected \( p < 0.001 \)), perturbation steps (Bonferroni corrected \( p < 0.001 \)), first recovery steps (Bonferroni corrected \( p = 0.03 \)) and second recovery steps (Bonferroni corrected \( p = 0.002 \)) for PC1. The difference in included angles was also significantly different from 5% asymmetry for PC2 when walking with 15% asymmetry during baseline right steps (Bonferroni corrected \( p = 0.01 \)) and perturbation steps (Bonferroni corrected \( p = 0.003 \)) and second recovery steps (Bonferroni corrected \( p = 0.04 \)). Lastly, there was only an effect of the direction of asymmetry for PC2 (\( F(1, 79), p = 0.049 \)) during the baseline right step (Baseline 1).

4. Discussion

We investigated how step length asymmetry affected intersegmental coordination patterns during responses to treadmill-based slip perturbations during walking. Our central finding was that intersegmental coordination patterns observed during asymmetrical walking differed from symmetrical walking during both unperturbed walking and perturbation recovery. When combined with previous observations that the reactive control of overall WBAM is not influenced by asymmetry [7], these results indicate that healthy people use a flexible combination of intersegmental coordination patterns rather than invariant reactions to maintain WBAM during perturbation responses when walking with asymmetric gait patterns.

Variations in coordination patterns during asymmetrical walking likely resulted from changes in the momentum generated by the lower extremities to reach the target asymmetry. Since the distal segments of the lower limbs are relatively far from the body’s center of mass and have a high velocity, they make the largest contribution to changes in intersegmental coordination patterns. For example, to achieve a positive asymmetry, participants placed their left...
foot further in front of the center of mass and increased the extension of their right hip so that the right foot was further behind their COM at heel strike. To achieve this objective,
Fig 5. Included angle between PCs extracted during asymmetrical walking (5%, 10%, and 15%) and symmetrical walking for each step (\(p<0.001\), \(p<0.01\), \(p<0.05\)) for (A) PC1 and (B) PC2. (C) Integrated whole-body angular momentum during asymmetrical walking relative to symmetrical walking for each step. Blue: Right step; Pink: Left step; Filled box plots: PC1; Non-filled box plots: PC2. The shaded gray area indicated the standard deviation of permuted included angle for each step, and the black line indicated the mean of the distribution.

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participants had to increase swing velocity. This likely explains why we observed increased weights of the left foot as SLA increased during right steps in the first principal component since positive step length asymmetries required longer left steps and faster foot swing.

The observation that reactive control of WBAM is consistent across levels of asymmetry [7] despite the variation in intersegmental coordination observed here may indicate that WBAM is a task-level variable that is stabilized by the nervous system during perturbation recovery. This is consistent with the framework proposed by the uncontrolled manifold hypothesis (UCM), which argues that the central nervous system allows for variability over a manifold of solutions that all successfully stabilize a higher-level performance variable [33]. Here, WBAM would serve as a high-level performance variable that is stabilized through covariation of elemental, segmental-level momenta. For example, Papi et al. demonstrated a similar concept when they found no differences between people post-stroke and healthy individuals in COM displacement during the stance phase of walking despite between-group differences in lower extremity joint kinematics [34]. Therefore, it is possible that when dynamic stability is challenged during walking, the central nervous system carefully regulates WBAM while allowing variance in lower-level, intersegmental coordination patterns.

Table 2. Statistical results from the ANOVA examining the effects of asymmetry and direction on the included angle for each step type.

| Step Type | PC     | Factor     | numDF | denDF | F-value  | P value  |
|-----------|--------|------------|-------|-------|----------|----------|
| Baseline1 | PC1    | Asym       | 2     | 73    | 9.7      | <0.001   |
|           |        | Direction  | 1     | 77    | 2.4      | 0.13     |
|           |        | Asym:Direction | 2     | 74    | 0.6      | 0.55     |
|           | PC2    | Asym       | 2     | 72    | 14.4     | <0.001   |
|           |        | Direction  | 1     | 78    | 4.0      | 0.049    |
|           |        | Asym:Direction | 2     | 74    | 0.08     | 0.92     |
| Baseline2 | PC1    | Asym       | 2     | 72    | 5.7      | 0.005    |
|           |        | Direction  | 1     | 75    | 1.3      | 0.26     |
|           |        | Asym:Direction | 2     | 73    | 0.1      | 0.88     |
|           | PC2    | Asym       | 2     | 71    | 11.0     | <0.001   |
|           |        | Direction  | 1     | 74    | 0.007    | 0.93     |
|           |        | Asym:Direction | 2     | 72    | 2.2      | 0.12     |
| Perturbation | PC1  | Asym       | 2     | 73    | 19.0     | <0.001   |
|           |        | Direction  | 1     | 75    | 1.9      | 0.18     |
|           |        | Asym:Direction | 2     | 73    | 0.5      | 0.59     |
|           | PC2    | Asym       | 2     | 72    | 8.7      | <0.001   |
|           |        | Direction  | 1     | 73    | 1.3      | 0.25     |
|           |        | Asym:Direction | 2     | 72    | 0.68     | 0.51     |
| Recovery1 | PC1    | Asym       | 2     | 74    | 11.2     | <0.001   |
|           |        | Direction  | 1     | 78    | 0.1      | 0.74     |
|           |        | Asym:Direction | 2     | 75    | 1.3      | 0.29     |
|           | PC2    | Asym       | 2     | 72    | 9.1      | <0.001   |
|           |        | Direction  | 1     | 75    | 0.1      | 0.72     |
|           |        | Asym:Direction | 2     | 73    | 0.8      | 0.45     |
| Recovery2 | PC1    | Asym       | 2     | 72    | 8.7      | <0.001   |
|           |        | Direction  | 1     | 75    | 1.8      | 0.18     |
|           |        | Asym:Direction | 2     | 73    | 0.4      | 0.67     |
|           | PC2    | Asym       | 2     | 73    | 8.5      | <0.001   |
|           |        | Direction  | 1     | 77    | 1.9      | 0.18     |
|           |        | Asym:Direction | 2     | 74    | 0.2      | 0.84     |

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In this study, we provided visual information about the desired and actual step lengths at each foot-strike throughout all trials, including the perturbation and recovery steps. Participants were encouraged to achieve the target step lengths for as many steps as possible, and therefore participants may have relied on this feedback during perturbation recovery to return to their pre-perturbation walking patterns faster than they otherwise would without visual feedback. However, participants’ reactive response is unlikely to influence measures of momentum until late into the first recovery step as the step length information was only shown after the foot-strike of the first recovery step. It remains to be seen if patterns of inter-limb coordination would differ in the presence of asymmetries that are not guided by online visual feedback.

Although the reactive intersegmental coordination patterns were significantly different from those observed during unperturbed locomotion, the overall patterns were qualitatively similar across steps. Taken together, these results may reflect two keys aspects of coordination during perturbed walking. First, the qualitative similarity between pre- and post-perturbation patterns observed here and in previous work [19] may reflect the dominant coordination patterns that characterize both unperturbed and perturbed bipedal walking. In contrast, the statistical differences between pre- and post-perturbation coordination patterns may reflect the changes in coordination necessary to maintain balance in response to perturbations. Patterns of intersegmental coordination observed during responses to external perturbations during walking likely capture a combination of passive limb dynamics, stereotypical pattern generation, and reactive balance control responses [35].

We observed that the upper limbs’ contribution to the control of angular momentum in the sagittal plane was negligible compared with lower limb segments during perturbation recovery. Since a stepping response is sufficient to restore balance from the treadmill accelerations used in this study, increases in momentum from the lower extremities may have been sufficient to restore sagittal plane WBAM. Consistent with our findings, Pijnappels et al. also found that arm movements had a small effect on body rotation in the sagittal plane during tripping over obstacles which elicits excessive forward rotation similar to the current study [36]. However, during larger perturbations that trigger backward falls, the arms elevate to shift the body’s center of mass back within the base of support [4]. This difference in the role of the arms across studies of perturbation recovery may result from the use of a larger velocity and displacement of the foot in the Marigold et al. [4] study. However, it remains to be seen how systematic variation of the magnitude and direction of external perturbations influences the role of the upper extremities during balance recovery.

Our results may also have implications for understanding the potential effects of interventions designed to reduce gait asymmetries in people post-stroke, as this is a common rehabilitation objective in this population [37]. The data from the current study illustrate how the intact neuromotor system modulates coordination between the upper and lower extremities in response to perturbations when walking asymmetrically. Based on the current results, we would hypothesize that reducing asymmetry in people post-stroke would also affect their reactive control of interlimb coordination during perturbation responses. However, in addition to spatiotemporal and kinematic asymmetries, people post-stroke have impaired reactive control of balance due to transmission delays [38], muscle weakness [39] and decreased propulsion in the paretic leg [40]. These deficits may prevent the paretic leg from initiating a successful stepping response to prevent excessive body rotation following external perturbations. Further work is necessary to determine if reducing asymmetry in people post-stroke would affect their interlimb coordination during reactions to perturbations and impact their ability to restore WBAM during perturbation recovery.
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