How Older Adults Regulate Lateral Stepping on Narrowing Walking Paths

Meghan E. Kazanski [a], Joseph P. Cusumano [b], Jonathan B. Dingwell [a,*]

[a] Department of Kinesiology, The Pennsylvania State University, University Park, PA 16802 USA
[b] Department of Engineering Science & Mechanics, The Pennsylvania State University, University Park, PA 16802 USA

* Please address correspondence to:

Jonathan B. Dingwell   Email: dingwell@psu.edu

CRediT Author Statement

Meghan E. Kazanski: Conceptualization, Data Curation, Formal Analysis, Investigation, Methodology, Software, Validation, Visualization, Writing – Original Draft, Writing – Review & Editing. **Joseph P. Cusumano:** Conceptualization, Funding Acquisition, Methodology, Supervision, Writing – Review & Editing. **Jonathan B. Dingwell:** Conceptualization, Data Curation, Funding Acquisition, Methodology, Project Administration, Resources, Supervision, Validation, Writing – Review & Editing.

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ABSTRACT (245 words)

Walking humans often navigate complex, dynamically changing walking paths. To better understand falling risk, is critical that we first determine how older adults purposefully vary their steps in varying environments that challenge balance. Here, 20 young (21.7±2.6 yrs) and 18 older (71.6±6.0 yrs) healthy adults walked on a series of continuously-narrowing virtual paths. Participants could switch onto an "easier" path at any time. We applied our Goal Equivalent Manifold framework to quantify how healthy aging alters lateral stepping variability and underlying multi-objective regulation processes along these paths. We also computed participants' lateral stepping characteristics when they switched paths. All participants (but more so younger adults) reduced their lateral stepping variability to remain on the path. How participants varied their lateral steps as paths narrowed closely reflected their trading-off maintaining step width for lateral body position. Older adults exhibited consistent differences in the structure of their lateral stepping variability, which reflected their stronger regulation of step width compared to younger adults, presumably to afford balance. While older adults left the narrowing paths sooner, several of their lateral stepping characteristics remained similar to those of younger adults. While age did not degrade path walking performance *per se*, it did alter how participants balanced the competing stepping regulation processes that likely gave rise to age-altered lateral stepping variability. Thus, older adults used the flexibility afforded by multi-objective lateral stepping regulation in context-dependent ways to maintain highly adaptive stepping on complex, dynamically changing walking paths.

INTRODUCTION

Daily walking environments contain complex arrangements of natural and built features, including terrain irregularities (Matthis et al., 2018), and myriad fixed (Twardzik et al., 2019) and moving obstacles (Moussaïd et al., 2011). Such features define the often highly irregular walking paths individuals must navigate in common contexts like crowded sidewalks, or cluttered store aisles. Individuals must continuously enact goal-directed stepping adjustments to remain on these continuously changing paths. As a result, people rarely perform prolonged, straight-ahead walking in daily life (Orendurff et al., 2008).

Despite its limited generalizability to real-world contexts, steady walking on straight invariant paths remains a dominant paradigm for extracting locomotor control principles (Herold et al., 2019; Vistamehr et al., 2016; Wang and Srinivasan, 2014). A critical, but unaddressed, locomotor control problem with immense real-world relevancy is how people enact continuous stepping adjustments on complex, dynamically changing walking paths.

It is especially critical to study how older adults enact ongoing stepping adjustments. Older adults experience high rates of falls (Burns and Kakara, 2018) that often result from inadequate stepping adjustments in challenging environments (Li et al., 2006; Rubenstein, 2006). Active modulation of lateral (i.e., side-to-side) stepping is particularly crucial to maintaining balance (Bruijn and van Dieën, 2018) while walking (Dingwell and Cusumano, 2019) along irregular paths or while maneuvering (Desmet et al., 2022). Healthy older adults exhibit generally greater lateral stepping variability during continuous, straight-ahead walking (Owings and Grabiner, 2004; Skiadopoulos et al., 2020), although

connecting these changes to fall risk remains challenging (Beauchet et al., 2009; Brach et al., 2005; Moe-Nilssen and Helbostad, 2005). We should instead address how older adults *modulate* their lateral stepping on dynamically changing walking paths that better reflect the types of circumstances that are more likely precipitate falls.

Older adults indeed exhibit increased lateral stepping variability when traversing irregular walking paths, such as between intermittent obstacles (Maidan et al., 2018) or onto stepping targets (Sun et al., 2017; Zhang et al., 2021). The sources of such age-related increased variability remain unclear. Such age-related changes in lateral stepping variability may reflect detrimental increases in physiological noise (Christou, 2011) that reduce lateral stepping accuracy (Kuo and Donelan, 2010) and compromise locomotor stability (Roos and Dingwell, 2010). Alternatively, age-related altered variability may reflect beneficial neuromuscular compensation (Eckardt and Rosenblatt, 2018) that allows older adults to exhibit considerable variability in select movement variables (Todorov, 2004) while still reliably accomplishing the goal-directed walking (Dingwell et al., 2017) necessary in challenging environments (Kazanski et al., 2020). Before determining how age-related stepping variability relates to falling risk in older adults, we must first determine the extent to which lateral stepping variability arises from which underlying sources (e.g., noise, motor redundancy, or changing task goals), especially on dynamically changing walking paths that require purposeful, goal-directed stepping adjustments.

We previously identified how stepping variability arises from goal-directed stepping regulation processes using an analytical framework entailing Goal Equivalent Manifolds (GEMs) (Cusumano and Dingwell, 2013; Dingwell et al., 2010). The GEM framework is based on formulating *goal functions* that pose testable hypotheses on how humans regulate stepping movements to achieve different walking tasks (Dingwell et al., 2010). The GEM is the set of all combinations of step-to-step observables (e.g. foot placements) that equally achieve some hypothesized walking task goal like "maintain speed" (Dingwell et al., 2010) or "maintain step width" (Dingwell and Cusumano, 2019), etc. We previously used the GEM framework to demonstrate that versatile walking is facilitated by inherently *multi*-objective lateral stepping regulation: humans can trade-off between competing task goals to maintain either step width or lateral body position (Dingwell and Cusumano, 2019). During continuous, straight walking, healthy adults prioritize minimizing errors from a target step width, while to a much lesser extent correcting errors in their lateral body position from the center of their walking path (Dingwell and Cusumano, 2019). Humans adapt these lateral stepping regulation processes to different contexts by re-weighting prioritization of these step width vs. lateral body position goals (Desmet et al., 2022; Render et al., 2021) and/or increasing correction of errors in *both* variables (Dingwell et al., 2020; Kazanski et al., 2020). Healthy aging does *not* alter lateral stepping regulation during straight or laterally-perturbed walking (Kazanski et al., 2020). We expect that these same types of GEM analyses should allow us to discern how age affects the lateral stepping regulation that people enact when walking on dynamically changing paths.

Here, we presented healthy young and older adults with systematically-narrowing virtual walking paths. This paradigm presented participants with situations that we expected would result in them continuously adjusting their lateral stepping regulation to maintain both step widths *and* lateral body positions on these defined paths. We predicted all (young and older) participants would reduce their variability as

paths narrowed and would progressively increase emphasis given to regulating body position, trading-off step width regulation as paths narrowed. With respect to age, we expected the older adults would exhibit greater variability (Lowry et al., 2017) than the younger, but would also retain greater prioritization of step width to promote lateral balance. Noting that some older adults decline to engage with sufficiently narrow paths at all (Butler et al., 2015), we gave participants the option to leave the narrowing path at any time. We evaluated whether group differences in when participants left the narrowing path reflected age-dependent limitations in lateral stepping regulation.

METHODS (1,452 words)

Participants

We analyzed data from the same cohort of 20 young healthy (YH) and 18 older healthy (OH) adults as in (Kazanski and Dingwell, 2021) (Table 1). We consented and screened all participants prior to study participation using protocols approved by the IRB at Pennsylvania State University.

[Insert Table 1 about here]

Protocol and Data Collection

Participants performed five, 4-minute trials walking on a series of virtual paths projected onto a 1.2m wide treadmill (Motek M-Gait system, Amsterdam, Netherlands), as detailed in (Kazanski and Dingwell, 2021). Each trial consisted of eight path walking bouts. Each bout presented two parallel, 25m-long virtual paths projected onto the left and right treadmill belts (Fig. 1A). One path gradually narrowed from an initial width of 0.45m to a final width of 0.05m according to:

$$W_P(x) = 0.45m - (0.016 \cdot x) \tag{1}$$

where $W_P(x)$ is the narrowing path width and x is the walker's path progression, which increased from 0m to 25m during each bout. Meanwhile, the adjacent path on the opposite treadmill belt remained fixed at 0.45m wide. A new walking bout began every 25m, at which point the narrowing and fixed-width paths alternated sides of the treadmill (Fig. 1B). Treadmill speed was fixed at 0.75 m/s, with optic flow of the virtual environment matched.

[Insert Figure 1 about here]

As people walk along *any* path, they adjust each foot placement to trade off maintaining step width (to stay balanced) for lateral position (to stay on their path) (see Supplement #1) (Dingwell and Cusumano, 2019; Kazanski et al., 2020). Thus, as paths presented here narrowed, this task slowly but increasingly constrained where people could place their feet to negotiate this trade-off. This task directly challenged each participant's ability to balance those competing task goals of maintaining balance while staying on their path.

Participants began each bout on the narrowing path and were instructed to remain on it as long as possible, while minimizing stepping errors off of the path. Participants chose when to switch over to the adjacent fixed-width path. Participants completed a 60s wash-out trial of steady-state walking after each path walking trial. Participants rested ($\geq$ 2-min) between all trials.

Kinematic data were collected as described in (Kazanski and Dingwell, 2021). Briefly, participants wore 16 retroreflective markers. Only left and right heel and posterior pelvis markers were used for the present analyses. Marker trajectories were collected at 120 Hz using a 10-camera motion capture system (Oxford Metrics, Oxford, UK), then post-processed in Vicon Nexus and exported to MATLAB (MathWorks, Natick, MA).

Relevant Stepping Variables
As we walk, we must choose, at each new step, where to place each foot relative to our path. (Dingwell and Cusumano, 2019). Here, we designate $\{z_{Ln}, z_{Rn}\}$ to indicate the lateral (z-direction) left and right foot placements relative to the path center (Fig. 2A) at each step, n.

[Insert Figure 2 about here]

We previously showed (Dingwell and Cusumano, 2019) that when humans walk on paths with constant width, they do *not* directly regulate $\{z_{Ln}, z_{Rn}\}$. Instead, they coordinate $\{z_{Ln}, z_{Rn}\}$ to multi-objectively maintain some combination of step width (w_n) and lateral body position (z_{Bn}) relative to their path. Foot placements, $\{z_{Ln}, z_{Rn}\}$, are directly mathematically related to these goal states, $\{z_{Bn}, w_n\}$, as described in Supplement #1.

For each walking bout, we extracted step-to-step time series of $\{z_{Ln}, z_{Rn}\}$ using the lateral positions of respective left and right heel markers at each consecutive step $n \in \{1, \cdots, N\}$ (Zeni et al., 2008). We then computed time series of $\{z_{Bn}, w_n\}$ from these $\{z_{Ln}, z_{Rn}\}$ (see Supplement #1). We then also extracted corresponding step-to-step time series of $W_P(x_n)$ for each walking bout.

We analyzed only steps taken along the narrowing path, until the second-to-last step before path switching. Each path switch instance occurred when the pelvic centroid crossed the treadmill midline. We extracted the narrowing path width ($(W_P)_{switch}$) at each path switch instance.

We first pooled all time series for all subjects by group (YH or OH). Then, we separated these stepping data into overlapping "bins" that covered the range of gradually-decreasing path width (W_P), as in (Kazanski and Dingwell, 2021). We required at least two *consecutive* steps from any given trial in each bin; single steps were discarded. Sixty-six total overlapping bins each contained all steps that occurred over a 0.075m range of W_P, with new bins assigned every 0.005m of W_P. Each bin's designated width, W_P, was taken as the mid-point of possible W_P for that bin. This allowed us to characterize how lateral stepping changed along the narrowing path (e.g., Fig. 3A). For these pooled group data within each of the 66 bins, we then sub-sampled 50 data sets by extracting a random 95% of the data in that bin.

[Insert Figure 3 about here]

Stepping Variability
To characterize each group's stepping behavior within each bin, we first visualize how steps $\{z_{Ln}, z_{Rn}\}$ directly relate to goal states $\{z_{Bn}, w_n\}$ by plotting them in the $[z_L, z_R]$ plane (Fig. 3B) (Desmet et al., 2022; Dingwell and Cusumano, 2019). In this $[z_L, z_R]$ plane, orthogonal diagonal lines define Goal

Equivalent Manifolds (Cusumano and Cesari, 2006) for maintaining either constant z_B or constant w (see Supplement #1). The distribution of lateral stepping variability within the $[z_L, z_R]$ plane will this reflect how strongly people prioritize z_B vs. w regulation (Supplement #1).

We first plotted all group-wise lateral steps in $[z_L, z_R]$ for each sub-sample. We fit 95% prediction ellipses to these stepping distributions, defined by scaling the eigenvalues of the $\{z_L, z_R\}$ covariance matrices by the 95th percentile critical value of the Chi-Square Distribution ($\chi_2^2(0.05) = 0.5991$) (Desmet et al., 2022). Each such ellipse can then be fully defined by its area, aspects ratio (a measure of "shape"), and orientation (Supplement #1; Fig. S1-1).

We computed the area (RA) of each variability ellipse relative to its square path area:

$$RA = \frac{\pi \cdot [\chi_2^2(0.05)] \cdot \sqrt{\lambda_1 \lambda_2}}{W_P^2}. \tag{2}$$

A larger RA indicates participants varied their lateral steps over a greater proportion of their available path width, W_P. We computed the aspect ratio of each ellipse as the ratio of covariance matrix eigenvalues (λ_1/λ_2), where $\lambda_1/\lambda_2 \gg 1$ indicates strong elongation along the direction of the ellipse major axis. We computed the orientation ($\Delta\theta$) of each ellipse as the angular deviation of the ellipse major axis from the $+w^*$ GEM (Supplement #1; Fig. S1-1).

We then averaged all 50 sub-sample values of each feature within each bin. We plotted these average RA, λ_1/λ_2, and $\Delta\theta$ across W_P for each group: YH and OH. We expected both groups would (necessarily) increase their RA as W_P decreased. However, we expected OH would exhibit consistently more-variable (i.e., larger RA) steps across the range of W_P. We further expected both groups would increase their prioritization of z_B over w regulation as W_P narrowed, yielding lower λ_1/λ_2 and/or higher $|\Delta\theta|$ with decreasing W_P. However, we expected OH would exhibit consistently greater prioritization of w (thus higher λ_1/λ_2, lower $|\Delta\theta|$) relative to YH across the range of W_P.

Error Correction
The above measures do not quantify the extent to which walking people correct errors in w and/or z_B from each step to the next (Dingwell and Cusumano, 2015, 2019). Participants attempting to achieve some target w^* at each step will exhibit small deviations on any n^{th} step: $w_n' = w_n - w^*$. Such deviations might then be corrected on the next step: $\Delta w_{n+1} = w_{n+1} - w_n$. For each sub-sample in each bin, we plotted all combinations of Δw_{n+1} vs. w_n' (Fig. 3C). We computed linear slopes $M(w)$ of each plot to quantify the extent of error correction, where $M(w) = -1$ indicates "perfect" correction, $M(w) < -1$ over-correction, and $M(w) > -1$ under-correction (Dingwell and Cusumano, 2015). We used this same procedure to quantify z_B error correction: $M(z_B)$. We averaged all 50 sub-sample values of each $M(z_B)$ and $M(w)$ within each bin and plotted average $M(z_B)$ and $M(w)$ across W_P for each group.

We expected that as W_P decreased, both groups would increase correction of z_B (i.e., $M(z_B) \to -1$) by trading off correction of w (i.e., $M(w) \to 0$). We further expected that, relative to YH, OH would demonstrate stronger prioritization of w (i.e., $M(w)_{OH} < M(w)_{YH}$) and weaker prioritization of z_B (i.e., $M(z_B)_{OH} > M(z_B)_{YH}$).

When People Switched Paths

We also characterized each participant's lateral stepping at the mean instants they switched off of the narrowing path. We selected the group-wise bin nearest to each participant's average $(W_P)_{switch}$. We then used the group-average stepping variability $\{(RA)_{switch}, (\lambda_1/\lambda_2)_{switch}, (\Delta\theta)_{switch}\}$ and error correction $\{M(z_B)_{switch}, M(w)_{switch}\}$ at that bin to represent that participant's lateral stepping at their average path switch. We compared these statistically in Minitab (Minitab, Inc., State College, PA) using two-sample t-tests or Mann-Whitney U tests to assess age group differences for each measure. We used these comparisons to determine differences in OH lateral stepping regulation might explain why they switched off the narrowing path sooner, at wider $(W_P)_{switch}$ (Kazanski and Dingwell, 2021).

RESULTS (298 words)

Most OH adults switched walking paths sooner at wider path widths, $(W_P)_{switch}$, than YH (Fig. 4) (Kazanski and Dingwell, 2021).

[Insert Figure 4 about here]

As paths narrowed, both groups remained within the path bounds ($RA < 1$), but progressively varied their lateral steps across more of the available W_P (Fig. 5A, left) (see also Supplement #2). OH consistently varied their lateral steps over a greater proportion of W_P (i.e., larger RA) than YH.

Initially at wider W_P, both YH and OH variability was strongly aligned with the w^* GEM ($\lambda_1/\lambda_2 \gg 1$ and $\Delta\theta \approx 0°$; Fig. 5B-C, left), consistent with straight-ahead walking (Dingwell and Cusumano, 2019). As paths narrowed, both groups progressively reduced alignment with the w^* GEM *toward* alignment with the z_B^* GEM. This was indicated by decreasing $\Delta\theta$ and by λ_1/λ_2 that initially decreased, but then increased as $\Delta\theta$ decreased further. Relative to YH, OH adults' lateral stepping variability deviated less from the w^* GEM (Fig. 5B-C, left).

[Insert Figure 5 about here]

Initially at wider W_P, both YH and OH groups exhibited stronger correction of step width errors ($M(w) \rightarrow -1$) and weaker correction of position errors ($M(z_B) \rightarrow 0$; Fig. 6A-B, left), also consistent with straight-ahead walking (Dingwell and Cusumano, 2019). As paths narrowed, both groups decreased w error correction (less negative $M(w)$) while increasing z_B error correction (more negative $M(z_B)$). Relative to YH, OH generally prioritized w correction more ($M(w)_{OH} < M(w)_{YH}$) and z_B correction less ($M(z_B)_{OH} > M(z_B)_{YH}$) throughout (Fig. 6A-B, left).

[Insert Figure 6 about here]

At their respective $(W_P)_{switch}$, YH and OH participants exhibited similar $(RA)_{switch}$ (Fig. 5A, right) and $(\lambda_1/\lambda_2)_{switch}$ (Fig. 5B, right), but significantly smaller (closer to 0°) $(\Delta\theta)_{switch}$ (Fig. 5C, right). Relative to YH, when OH switched off of the narrowing path, they prioritized z_B correction ($M(z_B)_{switch}$) similarly (Fig. 6A, right), but w correction ($M(w)_{switch}$) more (Fig. 6B, right).

DISCUSSION (766 words)

Humans must continuously navigate walking paths defined by features of their daily environments (Matthis et al., 2018; Twardzik et al., 2019). They do this by exploiting the multi-objective nature of lateral stepping regulation (Dingwell and Cusumano, 2019) to carry out goal-directed stepping adjustments across multiple challenging contexts (Desmet et al., 2022; Dingwell et al., 2020; Kazanski et al., 2020; Render et al., 2021). Any age-related changes in ability to regulate stepping may become particularly relevant when older adults walk in challenging environments. Here, young and older adults walked on continuously-narrowing virtual paths. We applied our GEM framework for lateral stepping to quantify how healthy aging alters the structure of lateral stepping variability and stepping regulation processes on these dynamically changing walking paths.

Prior studies examined how aging (Arvin et al., 2016) and redundancy (Rosenblatt et al., 2014) contribute to altered stepping variability on different paths of constant width. This present study differed: here, the path presented narrowed continuously and never imposed exact foot placements, step widths, or lateral body positions. Participants could thereby execute any of an infinite number of $\{z_L, z_R\}$ combinations to remain on the path. To remain on these paths, all participants progressively reduced their net amount of lateral stepping variability, while filling increasing portions of path width (Fig. 5A, left). Furthermore, both groups traded-off correcting w errors to more strongly prioritize correcting z_B errors (Fig. 6A-B left). In doing so, they thus executed steps that were distributed away from w^* GEM alignment toward the z_B^* GEM (Fig. 5B-C, left). Thus, both age groups progressively and substantially deviated from the goal-directed stepping characteristics observed during continuous, straight-ahead walking in a manner consistent with our theoretical framework (Dingwell and Cusumano, 2019). This present study further demonstrates how walking humans use the flexibility afforded by multi-objective regulation processes to achieve highly adaptive lateral stepping in diverse contexts (Desmet et al., 2022; Dingwell and Cusumano, 2019; Dingwell et al., 2020; Kazanski et al., 2020; Render et al., 2021).

Healthy aging did not degrade these older adults' capacities to keep foot placements within the path bounds although they did consistently exhibit greater relative variability within the path bounds compared to young adults (Fig. 5A, left). This is consistent with prior findings from fixed-width narrow paths that also showed older adults generally did not reduce lateral stepping variability to the same extent as younger adults (Arvin et al., 2016; Schrager et al., 2008).

This study is the first, to our knowledge, to quantify how healthy aging affects not only the amount and structure of lateral stepping variability, but also step-to-step regulation, in a continuously-varying context. Older adults differed in that they structured their lateral stepping to be consistently more closely aligned to the w^* GEM (Fig. 5B-C, left). This corresponded with their consistently stronger w correction (Fig. 6B, left) and weaker z_B correction (Fig. 6A, left) relative to young adults. This increased prioritization of w regulation is consistent with findings that older adults tend to rely on wider, more variable lateral foot placements to maintain balance while walking (Hollman et al., 2011; Owings and Grabiner, 2004). Thus, our work shows healthy older adults further appear to exploit multi-objective lateral stepping regulation to re-weight z_B vs. w stepping goals without compromising primary task performance.

The same older adults studied here tended to switch off the narrowing paths sooner and this was mediated by their somewhat lower self-perceived balance abilities (Kazanski and Dingwell, 2021). Here, we show that despite their earlier transitions, these older adults retained similar values for 3 of 5 lateral stepping variability and regulation characteristics at switching instances (Figs. 5-6, right). These complimentary analyses thus further illuminate how age-related changes in stepping regulation on these paths still allowed these older adults to achieve appropriate task performance.

Older adults often fall from improper or inadequate stepping adjustments, especially on walking paths that change in more challenging environments (Li et al., 2006; Rubenstein, 2006). Here, our study went beyond generic measures of age-related increases in variability magnitude (Brach et al., 2005) to also quantify both the structure of lateral stepping variability and how it connects to underlying multi-objective stepping regulation processes that affect walking balance. We showed that age-related changes in the structure of stepping variability on a continuously-narrowing path reflect step-to-step adaptations in how older adults regulate their movements to promote balance. Our novel implementation of GEM analyses to this continuous, distinctly non-steady state task demonstrates that walking humans flexibly adjust their multi-objective lateral stepping regulation in both age- and context-dependent ways to continuously adjust steps in real time in response to ongoing changes in challenging walking environments.

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Tables:

Table 1 – Young Healthy and Older Healthy physical characteristics and assessment scores. Functional balance ability was assessed using Timed Up and Go (TUG) and Four Square Step Test (FSST). Balance confidence was assessed using the Activities-Specific Balance Confidence (ABC) Scale and Iconographical Falls Efficacy Scale (I-FES). All values except Sex are given as Mean ± Standard Deviation. Statistical p-values are from two-sample t-test or Mann-Whitney U tests, as relevant.

Characteristic:	Young Healthy (YH):	Older Healthy (OH):	p-value
Sex [M/F]	9 / 11	5 / 13	
Age [yrs]	21.7 ± 2.6	71.6 ± 6.0	**< 0.001**
Body Height [m]	1.73 ± 0.08	1.70 ± 0.10	0.305
Body Mass [kg]	69.9 ± 12.1	70.0 ± 10.5	0.983
Assessment:			
TUG [s]	6.62 ± 0.75	8.20 ± 1.92	**0.004**
FSST [s]	7.63 ± 1.37	9.19 ± 1.53	**0.002**
ABC Score [/100 %]	96.8 ± 2.5	93.1 ± 5.5	**0.023**
I-FES [/40]	12.3 ± 1.9	16.1 ± 3.9	**0.001**

Figures:

Figure 1 – (A) For each walking bout, participants walked on a continuously-narrowing virtual path in an "M-Gait" system (Motek, Amsterdam, Netherlands). Participants were instructed to remain on the path as long as possible, while minimizing stepping errors outside of the path bounds. The narrowing path width gradually decreased from 0.45m to 0.05m. Meanwhile, an adjacent path remained fixed at 0.45m. Participants could switch off the narrowing path to the adjacent path at any time. **(B)** Each trial consisted of eight walking bouts. Narrowing vs. fixed-width paths alternated sides of the treadmill every 25m, at which point a new walking bout started and the path progression (x) was reset. Participants always switched to the fixed-width path before each new walking bout started. Example series of lateral (z-direction) left and right foot placements are shown as purple and green markers, respectively. Path switches (blue crosshairs) occurred when the lateral CoM trajectory crossed the centerline of the treadmill (dashed grey line). Dashed black lines indicate walking path centers.

A

B

Figure 2 – **(A)** Walking humans coordinate left and right foot placements from step-to-step to regulate lateral body position relative to path center and step width. Each step is defined by both foot placements (z_L, z_R) associated with that step. Lateral body position is the midpoint between them: $z_B = \frac{1}{2}(z_L + z_R)$. Step width is the difference between them: $w = (z_R - z_L)$ (Dingwell and Cusumano, 2019). **(B)** At any $W_P(x_n)$, successful left and right foot placements will land within the path bounds (Eq. 1), represented as a square region in the $[z_L, z_R]$ plane, centered at (0,0) with area $[W_P(x_n)]^2$. As $W_P(x_n)$ decreases along the path, the region of viable foot placements also decreases from wide (light grey square), to intermediate (darker grey square), to narrow (darkest grey square).

Figure 3 – **(A)** Lateral stepping time-series data are shown for a single OH participant. Each marker represents a single step at corresponding path width (W_P). Variables relative to path center include: left (z_L; purple) and right (z_R; green) foot placements (left), lateral body position (z_B; middle) and step width (w; right). For $\{z_L, z_R\}$ and z_B plots, grey dashed lines indicate path center. Two adjacent, overlapping bins are depicted in blue, then grey in the w plot. Each bin contains all foot placements within that bin's range of path widths. The first bin, centered at $W_P = 0.4125$m, contained all steps taken on path widths between 0.45m and 0.375m. The subsequent bin, centered at $W_P = 0.4075$m, contained all steps taken on path widths between 0.445m and 0.370m. For subsequent analyses, such lateral stepping time-series data were pooled separately for each YH and OH age groups across walking bouts, trials and participants. **(B)** Group-wise lateral foot placements plotted in $[z_L, z_R]$ at a given W_P. Individual steps are shown as single markers. Lateral stepping regulation variables $[z_B, w]$ map onto $[z_L, z_R]$ as mutually-orthogonal GEMs representing constant lateral body position ($z_B^* =$ constant; orange) and constant step width ($w^* =$ constant; green). The fitted 95% prediction ellipse (blue) is centered at the mean operating point (z_L^*, z_R^*) (black dot). The major ellipse axis is represented by the blue dashed line. **(C)** Group-wise $\{w'_n, \Delta w_{(n+1)}\}$ demonstrate deviations from w^* at step n ($w'_n = w_n - w^*$) and corrections on subsequent step $n+1$ ($\Delta w_{(n+1)} = w_{(n+1)} - w_n$) at a given W_P. Individual stepping combinations are represented by single markers. The diagonal solid blue line indicates the linear error correction, with slope $M(w)$ computed via least-squares fits that defines the extent of w correction in this sub-sample. The diagonal dashed black line indicates "perfect" error correction: i.e., $M(w) = -1$. These analyses were applied within all sub-samples across all quasi-constant W_P. Identical analyses were applied to group-wise $[z_B'_n, \Delta z_{B(n+1)}]$.

Figure 4 – Points along the narrowing paths at which younger (YH; left) and older (OH; right) participants transitioned off of the path. Shaded isosceles trapezoids represent the narrowing paths walked on. The left side axis labels give the distances traveled along the path (i.e., in the $+x$ direction). The right side axis labels give path widths as defined by Eq. (1). Individual average transition points are shown as circles for each participant (YH blue; OH red). Box plots indicate medians, 1st and 3rd quartiles, and whiskers extending to 1.5× interquartile range. OH participants switched walking paths sooner, at wider path widths (p = 0.022; Kazanski et al., 2021).

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Figure 5 – 95% prediction ellipses (Figs. 3B, 4) characterized lateral stepping variability in [z_L, z_R] across bins of decreasing path widths (W_P). **(A-C**, left**):** Relative area (RA), aspect ratio (λ_1/λ_2), and orientation ($\Delta\theta$), each averaged across all sub-samples within each bin (see Fig. 2). Shaded error regions represent ±1 SD between sub-samples. The path width bin at the average instant each participant switched paths, (W_P)$_{switch}$, was mapped onto each of these plots (shown in (**C**), left) to characterize their variability thresholds at path switch: i.e., (RA)$_{switch}$, (λ_1/λ_2)$_{switch}$, and ($\Delta\theta$)$_{switch}$. **(A-C**, right**):** OH exhibited similar (RA)$_{switch}$, (YH: 0.36±0.10; OH: 0.37×±0.20; p = 0.926) and (λ_1/λ_2)$_{switch}$ (YH: 1.26±0.03; OH: 1.27±0.07; p = 0.759), but less-negative ($\Delta\theta$)$_{switch}$ (YH: −117.38±13.33° ; OH: −61.12±28.93°; p = 1.54×10^{-7}) relative to YH.

Figure 6 – Average of sub-sampled linear error correction slopes for **(A**, left**):** lateral body position ($M(z_B)$) and **(B**, left**):** step width ($M(w)$ across bins of decreasing path widths (W_P). Shaded error regions represent ±1 SD between sub-samples. The path width bin at each participant's average instant of path switch, (W_P)$_{switch}$, was mapped onto each of these plots to determine error correction thresholds $M(z_B)_{switch}$ and $M(w)_{switch}$, as demonstrated in **(B**, left**)**. **(A-B**, right**):** OH exhibited similar $M(z_B)_{switch}$ (YH: −0.38±0.03; OH: −0.37±0.06; p = 0.737) and more-negative $M(w)_{switch}$ (YH: −0.39±0.03; OH: −0.56±0.03; p =2.33×10^{-19}) relative to YH.