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4 Quantifying template signatures of center-of-mass motion during walking with ankle 5 exoskeletons

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7 Michael C. Rosenberg^{1*}, Joshua L. Proctor^{1,2}, Katherine M. Steele¹

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11 ¹Department of Mechanical Engineering, University of Washington, Seattle, United States of America

12 ²Department of Applied Mathematics, University of Washington, Seattle, United States of America

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15 * Corresponding author

16 Email: mcrose3@emory.edu (MCR)

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18 Abstract

19 Predicting ankle exoskeleton impacts on an individual's walking function, stability, and efficiency
20 remains challenging. Characterizing how the dynamics underlying center-of-mass (COM) mechanics and
21 energetics change with exoskeletons may improve predictions of exoskeleton responses. We evaluated
22 changes in individual-specific COM dynamics in unimpaired adults and one individual with post-stroke
23 hemiparesis while walking in shoes-only and with passive ankle exoskeletons. We leveraged hybrid
24 sparse identification of nonlinear dynamics (Hybrid-SINDy) – an equation-free data-driven method for
25 inferring nonlinear hybrid dynamics using a large library of candidate functional forms – to identify
26 functional forms that best modelled physical mechanisms describing leg-specific COM dynamics, termed
27 *template signatures*. Across participants, Hybrid-SINDy identified template signatures comprised of leg
28 stiffness and resting length, similar to common spring-loaded inverted pendulum models. Rotary stiffness
29 mechanisms were identified in only 40-50% of participants. Unimpaired template signatures did not
30 change with ankle exoskeletons ($p > 0.13$). Conversely, post-stroke paretic leg and rotary stiffness
31 increased by 11% with zero- and high-stiffness exoskeleton, respectively, suggesting that COM dynamics
32 may be more sensitive to exoskeletons following neurological injury. Agreement between our
33 automatically-identified template signatures and those found from decades of biomechanics research
34 supports Hybrid-SINDy's potential to accelerate the discovery of mechanisms underlying impaired
35 locomotion and assistive device responses.

36

38 Introduction

39 Ankle exoskeletons are prescribed and designed to improve walking function and gait mechanics [1-3].
40 Personalized passive and powered ankle exoskeletons have shown promise to improve walking function
41 and muscle coordination in both unimpaired adults and individuals with neurological injuries [2, 4, 5].
42 However, changes in gait in response to ankle exoskeletons are highly individualized, especially
43 following neurological injury, making customization critical to improving device efficacy. For example,
44 in stroke survivors and children with cerebral palsy, ankle exoskeletons elicit diverse – and sometimes
45 detrimental – impacts on gait mechanics, walking speed, step length, and the energetic cost of walking[1-
46 4]. Capturing and characterizing these responses remains challenging and hinders clinicians and
47 designers’ abilities to customize exoskeletons to support walking function.

48

49 Despite observed changes in gait mechanics and energetics with ankle exoskeletons, little is known about
50 how task-level mechanisms, such as the control of whole-body dynamics, impact exoskeleton responses
51 [2, 5-8]. Specifically, controlling center-of-mass (COM) motion is known to impact gait energetics and
52 stability, and is related to clinical metrics of walking function and speed [9-11]. Modulating COM
53 dynamics in response to exoskeletons may, therefore, be critical to maintaining stable and efficient gait.
54 Importantly, following neurological injuries (*e.g.*, a stroke), COM energetics are altered asymmetrically,
55 with the paretic limb doing less work on the COM [12]. Similarly, when walking with ankle exoskeletons,
56 stroke survivors struggle to modulate their gait in energetically efficient ways [2]. However, these
57 experimental studies did not quantify changes in the underlying COM dynamics with exoskeletons. Even
58 in unimpaired adults it is unclear whether individuals modulate COM dynamics similarly in response to
59 exoskeletons. Characterizing the strategies used to modulate COM dynamics with ankle exoskeletons
60 may provide a foundational step towards understanding responses to ankle exoskeletons.

61

62 Reduced-order representations of gait dynamics provide a foundation to quantify and understand complex
63 exoskeleton responses [13-18]. For example, Full & Kodistchek (1999) used reduced-order
64 representations of COM dynamics, such as the spring-loaded inverted pendulum (SLIP), to quantify
65 strategies to achieve task-level goals including COM stabilization and perturbation responses during
66 locomotion [16]. Such reduced-order representations, or models, of gait encode neural and biomechanical
67 dynamics using a minimalist set of physics-based mechanisms. Specifically, common reduced-order
68 models of COM dynamics employ a variety of mechanisms, such as the SLIP leg springs or the rigid legs
69 of an inverted pendulum walker, describe COM accelerations during gait [10, 14, 16, 17, 19-23]. Each
70 mechanism within a model, therefore, encodes a hypothesis about how neural and biomechanical
71 subsystems interact to achieve task-level objectives. However

72

73 Reduced-order models of locomotion, often termed *Template* models, have provided influential insights
74 into strategies to control COM motion to achieve stable and efficient walking. For example, inverted
75 pendulum templates have been used to study COM energetics and the transition from walking to running
76 [17, 19], as well as strategies for energetically-efficient COM acceleration [9, 23, 24] and lateral
77 stabilization [10, 25]. Alternative template structures, such as the bipedal spring-loaded inverted
78 pendulum (SLIP) and its variants further improved predictions of sagittal-plane ground reaction forces
79 (GRFs) and revealed COM stabilization strategies across different speeds [14, 21, 22, 26, 27]. Extensions
80 of the bipedal inverted pendulum and SLIP templates include leg damping elements, rotary springs, or
81 curved feet. Note that each mechanism describes a unique relationship between leg kinematics and COM
82 acceleration [20, 28]. Further, given the breadth of templates proposed for human gait, the mechanisms
83 that best describe gait are likely individual-specific. Consequently, selecting an optimal template structure
84 to describe an individual's gait is non-trivial, particularly following neurological injury [29].

85

86 Template models that describe COM dynamics during unimpaired gait may be useful to serve as
87 normative archetypes, from which deviations can be used to characterize altered COM dynamics.

88 To emphasize the individualized nature of templates, we denote the subject-specific mechanisms that best
89 describe COM dynamics as an individual's *template signature*. For example, differences in the template
90 signatures in children with hemiparetic cerebral palsy differed from typically developing children and
91 between legs, with increased stiffness in the paretic limb [29, 30]. These differences suggested that
92 children with cerebral palsy may have reduced dynamic resources available to control COM motion
93 during walking, providing insight into individualized strategies to accelerate the COM and their
94 implications for walking function.

95

96 Similarly, we may expect that template signatures will reveal altered COM dynamics following other
97 forms of neurological injury (e.g., stroke) and with ankle exoskeletons. However, characterizing changes
98 in template signatures in these contexts requires addressing a major methodological challenge: Manually
99 identifying optimal template signatures mechanisms is a slow, *ad hoc* process that relies on first-
100 principles knowledge of the system [20]. The candidate template signature increases combinatorially with
101 the number of candidate mechanisms. Therefore, new methods are needed to select mechanisms rapidly
102 and systematically from a literature-based library of candidate mechanisms.

103

104 Recent advances in data-driven modeling and machine learning provide powerful techniques to flexibly
105 identify system dynamics from data [31-34]. **Hybrid-SINDy** (*SINDy: Sparse identification of nonlinear*
106 *dynamics* [31]) identifies sparse nonlinear dynamics in hybrid systems from time-series data, making it
107 particularly relevant to identifying template signatures of walking, which often have distinct dynamics
108 based on foot contact configuration [29, 33]. Importantly, Hybrid-SINDy identifies a large number of
109 candidate dynamical models (e.g., template signatures) from an arbitrary library of possible functional
110 forms (*i.e.*, mechanisms). The algorithm uses information criteria to determine levels of support for each
111 candidate model and selects only those that are parsimonious *and* highly representative of the system
112 [35]. Consequently, this approach enables rapid, systematic identification of individual-specific template
113 signatures while leveraging first-principles knowledge of the physics of human gait.

114

115 The purpose of this study was to identify changes in template signatures in response to ankle exoskeletons
116 and evaluate how neurological injuries may alter template signatures in response to exoskeletons. We
117 used the Hybrid-SINDy algorithm to identify template signatures that encode COM dynamics during
118 walking in unimpaired adults and how those signatures changed with hinged and stiff ankle exoskeletons.
119 We hypothesized that ankle exoskeletons would primarily increase leg stiffness. Additionally, to examine
120 the potential of template signatures to encode changes in COM dynamics for individuals with
121 neurological injuries, we present a case study evaluating altered template signatures in an individual with
122 post-stroke hemiparesis. To our knowledge, this work is the first to employ physics-informed data-driven
123 modeling techniques to select low-dimensional, interpretable, representations of walking dynamics with
124 ankle exoskeletons.

125 Methods

126 Data collection

127 We collected 3D marker trajectories using a ten-camera motion capture system (Qualisys AB,
128 Gothenburg, SE) and ground reaction forces (GRFs) using an instrumented treadmill (Bertec Inc.,
129 Columbus, USA) in twelve unimpaired adults (6M/6F; age = 23.9 ± 1.8 years; height = 1.69 ± 0.10 m;
130 mass = 66.5 ± 11.7 kg) and one stroke survivor with lower-limb hemiparesis (*sex not disclosed*; age = 24
131 years; height = 1.70 m; mass = 68.0 kg). Participants walked on a treadmill in shoes-only and with
132 bilateral passive ankle exoskeletons under two conditions: zero resistance to ankle flexion (*i.e.*, zero
133 stiffness; K_0) and high dorsiflexion resistance (*i.e.*, high stiffness; $K_H = 5.1$ Nm/deg; Figure 1). A detailed
134 description of the experimental protocol and data preprocessing can be found in [36]. Briefly, in a second
135 session following a practice session, data were collected while participants walked at their self-selected
136 speed for four minutes per condition. To mitigate fatigue, the post-stroke participant walked under the
137 same protocol, but for only three minutes per condition [37]. This study was approved by the University
138 of Washington Institutional Review Board (#47744) and all participants provided written informed
139 consent to participate in the study.

140

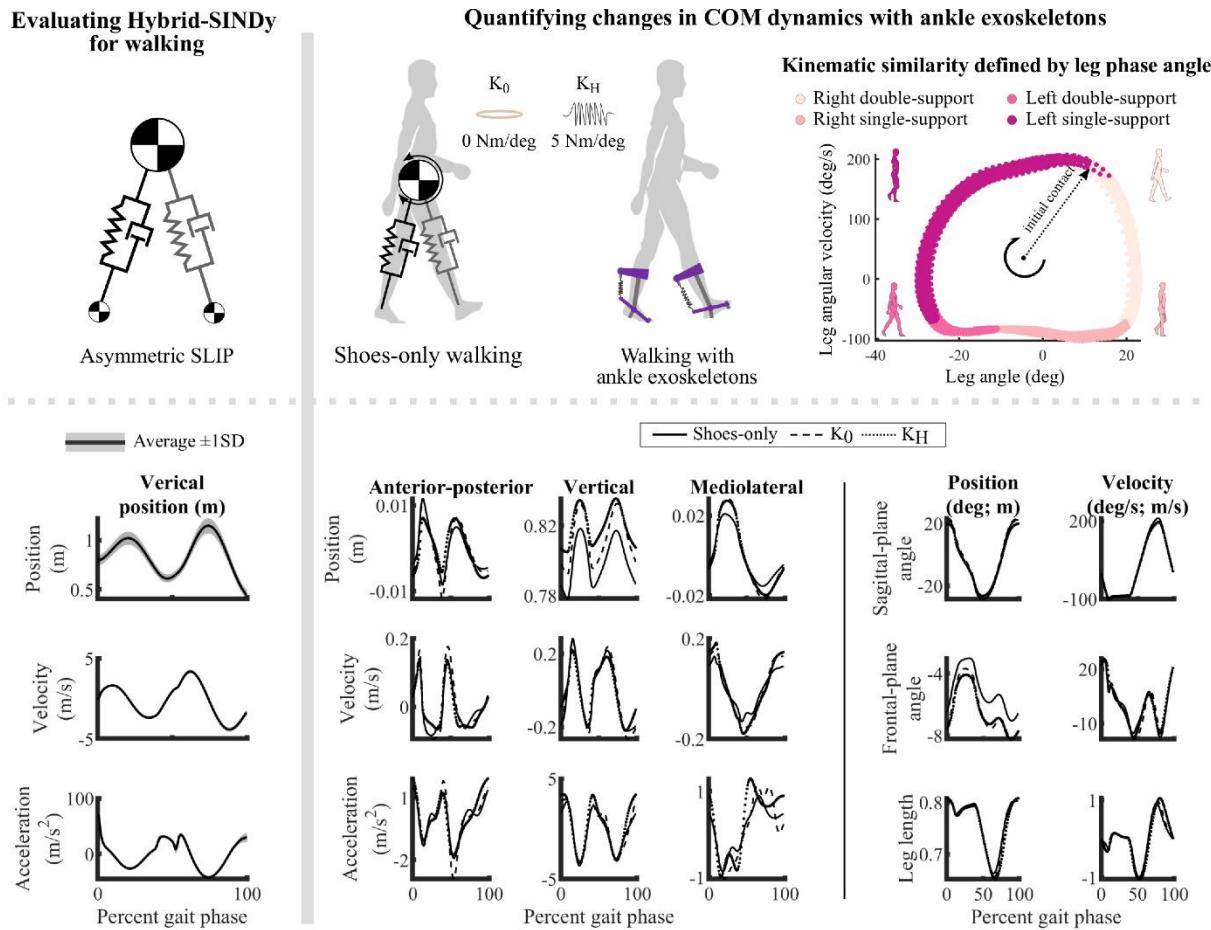


Figure 1 – **Top:** Two-dimensional depictions of the simulated SLIP (left) and human walking (right). The simulated SLIP used leg springs and dampers, as well as foot masses to simulate full strides. Participants walked on a treadmill in shoes-only and in ankle exoskeletons under zero- (K_0) and high-stiffness (K_H) conditions. The pink phase portrait was used to define a kinematic phase variable for clustering similar measurements. Colors denote gait phases corresponding to first and second double-limb support, single-limb support, and swing. **Bottom:** Time-series measurements of COM position, velocity, and acceleration for the simulated SLIP (left; average \pm 1SD) and an exemplary unimpaired adult participant (right). For human walking, 3D COM states (middle) and leg states (right) are shown for the three exoskeleton conditions. The leg length and velocity, and sagittal- and frontal-plane leg angles determined leg forcing, while COM position determined the direction of forcing.

141

142 **Estimating template signatures with Hybrid-SINDy**

143 To identify mechanisms describing COM acceleration, we used the Hybrid-SINDy algorithm to identify
 144 template signatures with and without ankle exoskeletons. Note that, for each exoskeleton condition, we
 145 used OpenSim's *Body Kinematics* algorithm to estimate the COM and foot positions [38, 39]. In this

146 section, we describe the SINDy and Hybrid-SINDy algorithms in the context identifying template
147 signatures, while more detailed explanations can be found in [31] and [33, 34]. Let 3D COM
148 accelerations, $\ddot{\mathbf{q}}(t) \in \mathbb{R}^{m \times n}$, be described by continuous-time nonlinear dynamics, $f(\mathbf{q}(t), \dot{\mathbf{q}}(t))$, where
149 m denotes the number of samples and $n=3$ output:

$$\frac{d^2}{dt^2} \mathbf{q}(t) = \ddot{\mathbf{q}}(t) = f(\mathbf{q}(t), \dot{\mathbf{q}}(t)), \quad 1$$

150 where time is denoted by $t \in \mathbb{R}^{m \times 1}$, and $\mathbf{q}(t)$ and $\dot{\mathbf{q}}(t)$ represent COM positions and velocities relative to
151 the feet, respectively, in $\mathbb{R}^{m \times n}$, in the anterior-posterior, vertical, and mediolateral directions. We assume
152 that only a small number of functional forms (*i.e.*, mechanisms) in $f(\mathbf{q}(t), \dot{\mathbf{q}}(t))$ describe most of the
153 system's behavior. We omit the time notation in the remaining sections.

154

155 Sparse Identification of Nonlinear Dynamics (SINDy)

156 The SINDy algorithm ([31]) recovers sparse nonlinear dynamics from a library of candidate functional
157 forms, which may consist of arbitrary nonlinear functions of system measurements. Adopting the notation
158 from [33], we can rewrite the dynamics in equation 1 as:

$$\ddot{\mathbf{q}} = \Theta(\mathbf{q}, \dot{\mathbf{q}})\Xi, \quad 2$$

159 where $\Xi \in \mathbb{R}^{p \times n}$, is a linear map from nonlinear function library encoding mechanisms potentially
160 describing COM accelerations, $\Theta(\mathbf{q}, \dot{\mathbf{q}}) \in \mathbb{R}^{m \times p}$, to COM accelerations, $\ddot{\mathbf{q}}$. The coefficients in Ξ ,
161 therefore, represent template signatures parameters, while Θ encodes template signature structure. We
162 included $p = 14$ functional forms in the function library, described below. The SINDy algorithm
163 promotes sparsity using sequential least-squares regression with hard thresholding, with the threshold
164 defined by the sparsity parameter, λ (Eq 3) [33, 34]. This thresholding approach penalizes the zero-norm
165 of Ξ and solves:

$$\min_{\Xi} \|\Theta(\mathbf{q}, \dot{\mathbf{q}})\Xi - \ddot{\mathbf{q}}\|_2 + \lambda \|\Xi\|_0, \quad 3$$

166

167 Hybrid-SINDy

168 Hybrid-SINDy extends SINDy in two important ways. First, Hybrid-SINDy uses clustering to generalize
169 SINDy to hybrid systems. For human walking data clustering, enables unique dynamics to be identified in
170 each hybrid regime, such as those defined by contact configuration (*i.e.*, single- and double-limb support)
171 [11, 15, 19]. Second, Hybrid-SINDy uses information criteria to automatically select the system dynamics
172 that best describe the data. This approach enables competing hypotheses about the mechanisms describing
173 COM acceleration to be rapidly and systematically compared, thereby highlighting mechanisms that are
174 common across individuals and those unique to a subset of individuals.

175

176 [Applying Hybrid-SINDy to walking](#)

177 We applied the Hybrid-SINDy algorithm to human gait using the following steps for each participant and
178 walking condition (Figure 2). Note that within each hybrid regime of the gait cycle, we expanded upon
179 the original Hybrid-SINDy algorithm by using multi-model inference to define a single template signature
180 when multiple signatures were plausible (Step 5) [33, 34].

- 181 1. *Clustering*: For each sample in the training set, we generated cluster the samples' 800 nearest
182 neighbors and identified the centroid of each cluster. The average cluster width (7.4% of a stride) was
183 less than double-limb support. We used the right leg angle and angular velocity with respect to
184 vertical as our clustering parameters [36].
- 185 2. *Model estimation*: For each cluster, we used SINDy to identify multiple template signatures, by
186 sweeping 40 sparsity threshold values, typically producing 5-15 unique signatures per cluster.
- 187 3. *Model evaluation and selection*: Using held-out data, we evaluated the ability of each template
188 signature to reconstruct COM accelerations in the anterior-posterior, vertical, and mediolateral
189 directions. We computed the average reconstruction error of held-out data over gait phases, ϕ ,
190 spanning 0-100% of a stride (Eq 4).

$$\text{error}_\phi = |\Theta(\mathbf{q}, \dot{\mathbf{q}}) \mathbf{\Xi} - \ddot{\mathbf{q}}|_\phi$$

191 We selected template signatures based on two criteria: First, we discarded signatures that were
192 identified in less than 1% of training clusters. Frequently occurring template signatures are more
193 likely to be robust to measurement noise or stride-to-stride variability, making them better
194 representations of an individual's gait dynamics [33].

195
196 Second, for each hybrid regime – single- and double-limb support – we selected the frequently
197 occurring template signatures that had the highest likelihood of being the correct signature describing
198 COM accelerations according to the Akaike Information Criterion (AIC) [35, 40]. The AIC is widely
199 used to compare candidate representations of system dynamics (*e.g.*, template signatures) according
200 to their number of free parameters and log-likelihood [35]. The AIC favors parsimonious, highly-
201 representative models, which is ideal for identifying minimalist representations of gait dynamics.
202 Like Mangan and colleagues, we used the AIC corrected for finite sample sizes (AICc) and compared
203 signatures according to their relative AICc score ($\Delta_j = AICc_j - AICc_{min}$ for signature j) [33, 35].
204 The best model according to the relative AICc score has a score of $\Delta_j = 0$ and all other models had
205 higher scores. Burnham and Anderson noted that models with $\Delta_j \leq 2$ have substantial support, while
206 $\Delta_j > 7$ have low support [35]. We adopted the threshold of [33], deeming template signatures with
207 $\Delta_j \leq 3$ to be *plausible*.

208 4. *Multi-model inference*: Since human gait dynamics are not strictly hybrid, multiple template
209 signatures are likely to be deemed plausible in each hybrid regime. To construct a single template
210 signature for each hybrid gait regime, we computed a weighted-average signature using Akaike
211 weights, ω_j , where j is the j^{th} plausible model in a hybrid regime [35]. Akaike weights are

$$\omega_j = \frac{\exp\left(-\frac{\Delta_j}{2}\right)}{\sum_{r=1}^R \exp\left(-\frac{\Delta_r}{2}\right)},$$

212

213 where $\exp\left(-\frac{4j}{2}\right)$ defines the likelihood of the j^{th} template signature given the observations [35]. This
 214 approach weights each signature based on its likelihood relative to the other plausible signatures.
 215 5. *Uncertainty estimation*: To evaluate the robustness of the template signatures to noise and stride-to-
 216 stride variations in the data, we performed 200 bootstrapped estimates of template signatures
 217 parameters using the data in the corresponding hybrid regimes. The template signatures were
 218 characterized by the mean and standard deviation of the bootstrapped parameters.
 219

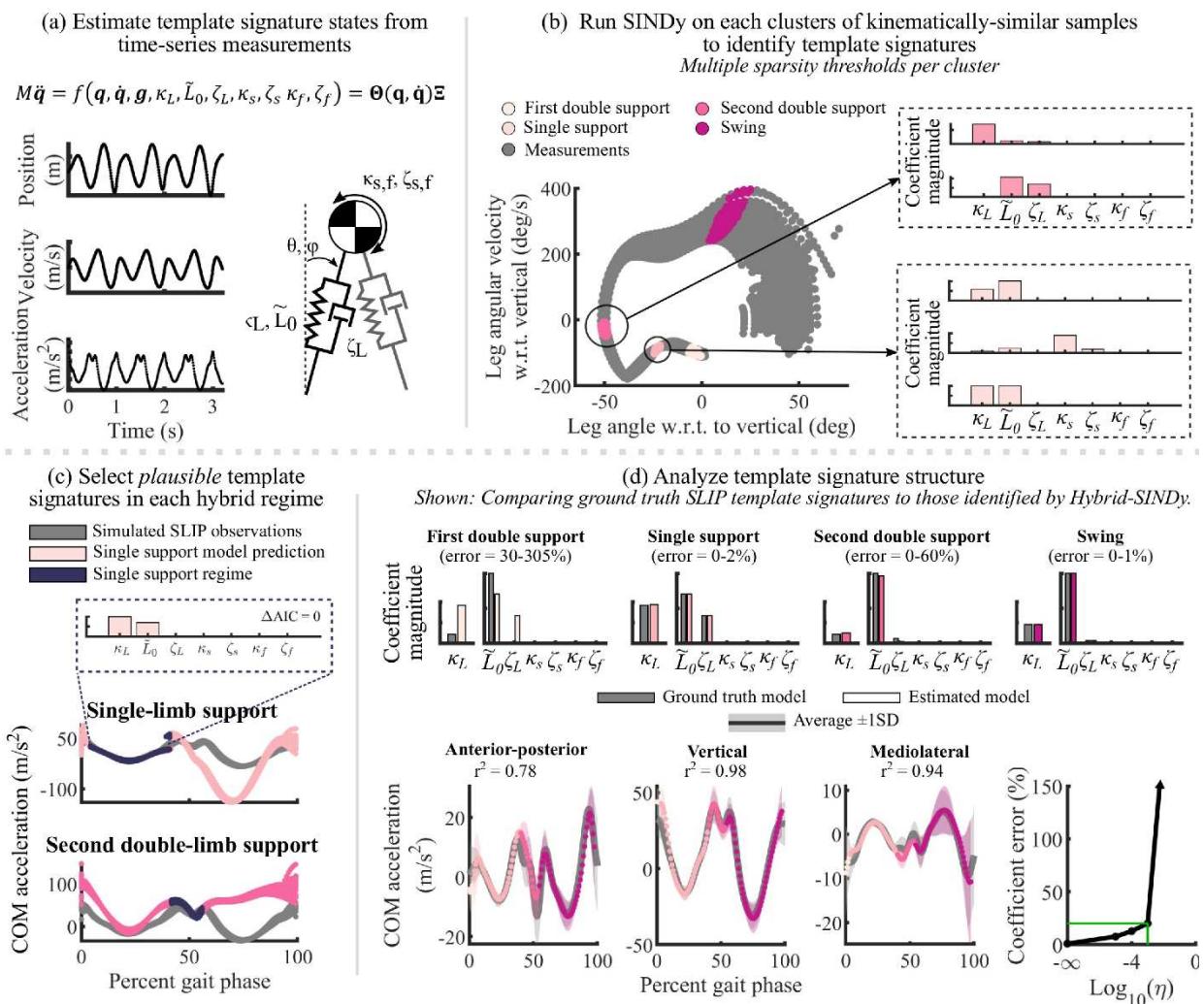


Figure 2 – The Hybrid-SINDy algorithm, applied to a simulated 3D-SLIP walking model. (a) COM and foot measurements from the simulated SLIP were used to approximate nonlinear SLIP states, $\Theta(\mathbf{q}, \dot{\mathbf{q}})$, as shown in the equation and the SLIP diagram. Variables can be found in Table 1. (b) Kinematically-similar measurements were used to cluster the data (gray) and exemplary template signature parameters from two clusters. For each cluster, multiple signatures of different complexities

were identified. (c) Each signature was evaluated in the hybrid regime containing the cluster centroid. The plots show COM accelerations and predictions in single-limb (top) and second double-limb (bottom) support. Error is low only in the correct hybrid regime for each signature (purple). (d) **Top:** Ground-truth (gray) and identified (colors) template signatures in each hybrid regime, with percent errors compared to ground-truth. **Bottom:** Average ($\pm 1\text{SD}$) observed (gray) and predicted (colors) COM accelerations. Colors denote each hybrid regime. The rightmost plot shows the average percent error in template signature coefficients during single-limb support and swing with increasing measurement noise (η). Green lines denote the approximate noise level of marker-based motion capture.

220

221 Template signatures mechanisms and dynamics

222 To model three-dimensional COM dynamics during walking, we created a function library of the
223 following candidate mechanisms based on prior literature (Table 1):

224 • *Rigid legs*, which reflected pendular gait in stance [10, 13, 15, 19, 24]. Pendular gait is energetically
225 passive and reflects kinetic and potential energy transfer during walking but cannot explain GRFs and
226 lacks double-limb support phase [15, 22, 41].

227 • *Leg springs* ([14, 20-22, 26]) and *dampers* ([28]), which produce force along the leg. Leg springs are
228 common energetically-conservative mechanisms used to describe walking and running dynamics, and
229 enable a double-limb support phase. Leg dampers are less common, but have been used to capture
230 non-conservative gait dynamics [28].

231 • *Rotary springs* ([14, 20]) and *dampers* in the sagittal and frontal planes, which enable forcing
232 transverse to the leg axis. The addition of a rotary springs has been show to improve reconstructions
233 of anterior-posterior GRFs in a bipedal SLIP [20]. We did not identify rotary damping elements in
234 prior literature but include them as a candidate mechanism describing non-conservative transverse
235 forcing.

236 The mechanisms selected by the Hybrid-SINDy algorithm define the template signature *structure*, which
237 describes characteristic strategies to accelerate the COM. The identified signature *parameters* describe
238 each mechanism's contribution to COM accelerations.

239

240 The dynamics of a three-dimensional bipedal SLIP augmented with damping and rotary mechanisms may
 241 be written as

$$M(\ddot{\mathbf{q}} - \mathbf{g}) = \sum_{j=R,L} \left(-[k_L(L - L_0) + c_L \dot{L}] \frac{\mathbf{q}}{L} - [k_s \theta + c_s \dot{\theta}] \frac{\mathbf{q}}{L_s^2} - [k_f \phi + c_f \dot{\phi}] \frac{\mathbf{q}}{L_f^2} \right)_j, \quad 6$$

242
 243 where M is body mass, \mathbf{g} is the gravity vector, ϕ describes the traverse-plane leg angle, and θ describes
 244 the leg angle from vertical in the direction defined by ϕ [20, 26]. The summation represents total force
 245 generated by the legs on the COM. The left-most brackets contain mechanisms that impart forces radially
 246 along the leg: k_L is the leg stiffness, L is the instantaneous leg length, L_0 is the leg resting length, c_L is the
 247 leg damping, \dot{L} is the instantaneous leg velocity. We henceforth denote L_0 as *leg length* for clarity. The
 248 middle bracket contains mechanisms that impart forces transverse to the leg axis in the sagittal plane: k_s
 249 and c_s are the sagittal-plane rotary stiffness and damping, respectively. L_s denotes the sagittal-plane leg
 250 projection. Analogously in the right-most brackets, k_f and c_f represent the frontal-plane rotary stiffness
 251 and damping, respectively, and L_f denotes the frontal-plane leg projection.

252
 253 **Normalized template mechanisms**
 254 To account for inter-individual differences in walking speed and body size during analysis, we
 255 normalized the template signatures during analysis (Table 1). Leg stiffness was normalized as in [18, 20,
 256 33]. Leg resting length normalized to the measured leg length [20, 28]. Rotary stiffness was normalized
 257 according to [20]. All damping terms were converted to damping ratios [28]. The normalized leg, sagittal-
 258 plane, and frontal-plane stiffness mechanisms are denoted by κ_L , κ_s , and κ_f , respectively. The normalized
 259 leg, sagittal-plane, and frontal-plane damping mechanisms are denoted by ζ_L , ζ_s , and ζ_f , respectively.
 260 Normalized leg length is denoted \tilde{L}_0 . We can rewrite equation 6 as a linear combination of our normalized
 261 coefficients and nonlinear transformations of our states:

$$M(\ddot{\mathbf{q}} - \mathbf{g}) = [\kappa_L \quad \kappa_L \tilde{L}_0 \quad \zeta_L \quad \kappa_s \quad \zeta_s \quad \kappa_f \quad \zeta_f]_j \Theta(\mathbf{q}, \dot{\mathbf{q}}), \quad j = \{R, L\} \quad 7$$

262

263 The COM position and velocity relative to the feet were used to compute candidate template signature
264 states: leg lengths and lengthening velocities, sagittal-plane leg angles and angular velocities relative to
265 vertical, and frontal-plane leg angles and angular velocities relative to vertical.

266

Table 1 – List of template signature terms.

Term	Symbol	Normalized form
Leg stiffness	κ_L	$\frac{k_L L_{bio}}{Mg}$
Leg resting length	\tilde{L}_0	$\frac{L_0/L_{bio}}{c_L}$
Leg damping	ζ_L	$\frac{c_L}{2\sqrt{k_L M}}$
Sagittal-plane rotary stiffness	κ_s	$\frac{k_s}{Mg L_{bio}}$
Sagittal-plane rotary damping	ζ_s	$\frac{c_s}{\sqrt{\kappa_s M L_{bio}^2}}$
Frontal-plane rotary stiffness	κ_f	$\frac{k_f}{Mg L_{bio}}$
Frontal-plane rotary damping	ζ_f	$\frac{c_f}{\sqrt{\kappa_f M L_{bio}^2}}$

c_L =leg damping; c_s =sagittal-plane rotary damping;
 c_f =sagittal-plane rotary damping; g = gravitational acceleration; k_L =leg stiffness; k_s =sagittal-plane rotary stiffness; k_f =sagittal-plane rotary stiffness;
 L_{bio} biological leg length; M = body mass

267

268 Evaluating Hybrid-SINDy’s ability to select template signatures

269 To evaluate Hybrid-SINDy’s ability to accurately identify and select walking dynamics, we evaluated the
270 algorithm on surrogate data of a simulated bipedal SLIP walker with known dynamics (Figure 1 & Figure
271 2) [26]. The bipedal SLIP had asymmetric leg stiffness and damping, and masses representing the pelvis
272 ($M = 56$ kg) and feet ($M_f = 7$ kg) to simulate a full gait cycle (Table 2). We omitted rotary mechanisms
273 from the SLIP dynamics but used the full function library described in equation 6. We simulated 120 gait
274 cycles from randomly perturbed initial conditions with an average initial velocity of 1.20 m/s. As

275 SINDy's performance is sensitive to sensor noise, we identified template signatures using simulation
276 measurements with added Gaussian noise ranging logarithmically from 10 nm to 10 cm applied to the
277 position measurements [31]. We quantified the Hybrid-SINDy's ability to reconstruct model dynamics
278 using the number of correctly-selected mechanisms and the accuracy of estimated coefficients.

279

Table 2 – Bipedal SLIP normalized simulation parameters.

Term	<i>[Right leg, Left leg]</i>			
	DS ₁	SS	DS ₂	SW
κ_L	[10.8, 9.1]	[25.1, 18.82]	[9.03, 10.68]	[36.6, 25.3]
\tilde{L}_0	[1.0, 1.0]	[1.0, 0.7]	[1.0, 1.0]	[0.7, 0.7]
ζ_L	[0.04, -0.07]	[-0.03, -0.04]	[0, 0]	[-0.40, -0.57]
κ_s	0	0	0	0
ζ_s	0	0	0	0
κ_f	0	0	0	0
ζ_f	0	0	0	0

280

281

282 Identifying template signatures in human gait

283 We estimated unique template signatures that predicted COM accelerations using the Hybrid-SINDy
284 algorithm for each participant and ankle exoskeleton condition. We used 10800 samples (90 seconds)
285 were available for clustering and we identified template signatures for the first 3600 samples. An
286 additional 3600 samples of held-out data were used for model evaluation and selection.

287

288 To evaluate the extent to which each mechanism described COM accelerations in unimpaired adults, we
289 analyzed the proportion of participants for whom each template signature coefficient was selected.

290 Signature terms are identified across individuals represent mechanisms fundamental to COM dynamics,
291 while infrequently identified mechanisms describe individual-specific features of COM dynamics.

292

293 To determine if unimpaired COM dynamics during shoes-only walking generalized to walking with ankle
294 exoskeletons, we evaluated the ability of shoe-walking template signature structures to reconstruct COM
295 accelerations in the K_0 and K_H conditions. We used least-squares regression to identify template signature
296 parameters for the K_0 and K_H conditions using the shoes-only template signature structure. We compared
297 the relative AICc ($\Delta AICc$) scores between these signature structures and those of the signature structures
298 specific to the K_0 and K_H trials. We used one-sample paired t-tests ($\alpha = 0.05$) to determine if shoes-only
299 template signatures were less plausible (*i.e.*, $\Delta AICc \geq 3$) than signature structures selected for the K_0 and
300 K_H conditions.

301

302 To determine the frame and mass of the ankle exoskeletons impacted COM dynamics, we compared
303 template signature coefficients between the K_0 and Shoe conditions. Similarly, to evaluate the impacts of
304 exoskeleton stiffness on COM dynamics, we compared template signature coefficients in the K_H and K_0
305 conditions. For both comparisons, we used paired two-sample t-tests with Holm-Sidak step-down
306 corrections for multiple comparisons ($\alpha = 0.05$) [42]. Because we found that shoes-only template
307 signature structures were plausible for most unimpaired participants, we used the shoes-only template
308 signatures structure to estimate signature coefficients in all exoskeleton conditions.

309

310 To determine if COM dynamics may be altered post-stroke, we first computed the percent difference in
311 the non-paretic and paretic limb template signature parameters during shoes-only walking in one
312 individual with post-stroke hemiparesis. We also evaluated changes in post-stroke COM dynamics with
313 ankle exoskeletons by computing percent changes in template signature coefficients for the K_0 condition
314 compared to the shoes-only and K_H conditions.

315

316

317 **Results**

318 **Hybrid-SINDy identified COM dynamics of a simulated SLIP**

319 The Hybrid-SINDy algorithm correctly identified the salient features of the simulated SLIP's template
320 signature during single-limb support (Figure 2). Specifically, Hybrid-SINDy identified non-zero leg
321 stiffness (κ_L), resting length (\tilde{L}), and damping (ζ_L) terms only – consistent with the SLIP's dynamics. In
322 the noise-free condition, 86% of terms were correctly selected and only one term, leg damping in first
323 double-limb support, was incorrectly selected. Both double-limb support regimes were very short in the
324 simulated SLIP: 1.3% and 4.6% of samples in first and second double-limb support – smaller than the
325 7.4% of the training data in each cluster. Consequently, all clusters with centroids in the double-limb
326 support regimes contained samples spanning double- and single-limb support. Hybrid-SINDy incorrectly
327 omitted only damping terms with small coefficients (Table 2). Without measurement noise, in single-limb
328 support and swing, coefficients were within 2% of the ground truth values. When measurement noise with
329 a standard deviation of 1 mm – similar to that of our motion capture system – was added to the system
330 measurements, Hybrid-SINDy correctly selected only 63% of mechanisms, though incorrect mechanisms
331 selection occurred primarily in double-limb support. Template signature coefficients were estimated with
332 an average error of less than 29% in single-limb support and swing (Figure 2). In single-limb support,
333 Hybrid-SINDy was still able to estimate large coefficients (*i.e.*, leg stiffness and resting length) with less
334 than 1% error.

335

336 **Shoes-only template signatures revealed common and subject-specific COM mechanisms**

337 When walking in shoes-only, SLIP-like template signatures best described COM dynamics across
338 unimpaired participants. In both double-limb and single-limb support, SLIP mechanisms – leg stiffness
339 and leg length – were selected in 96-100% of limbs (Figure 3). Unimpaired template signatures were not
340 significantly different between legs (paired 2-sample t-test; $p > 0.01$). However, template signatures
341 varied during double-limb support: Rotary stiffness terms were selected in 58-67% of limbs in the first

342 double-limb support and 83-95% of limbs during second double-limb support. Except for sagittal-plane
343 damping, damping terms were selected in less than 20% of limbs. As expected during swing, when the
344 limb should not contribute to COM accelerations, no terms were selected. We omit the swing regime from
345 remaining analyses.

346

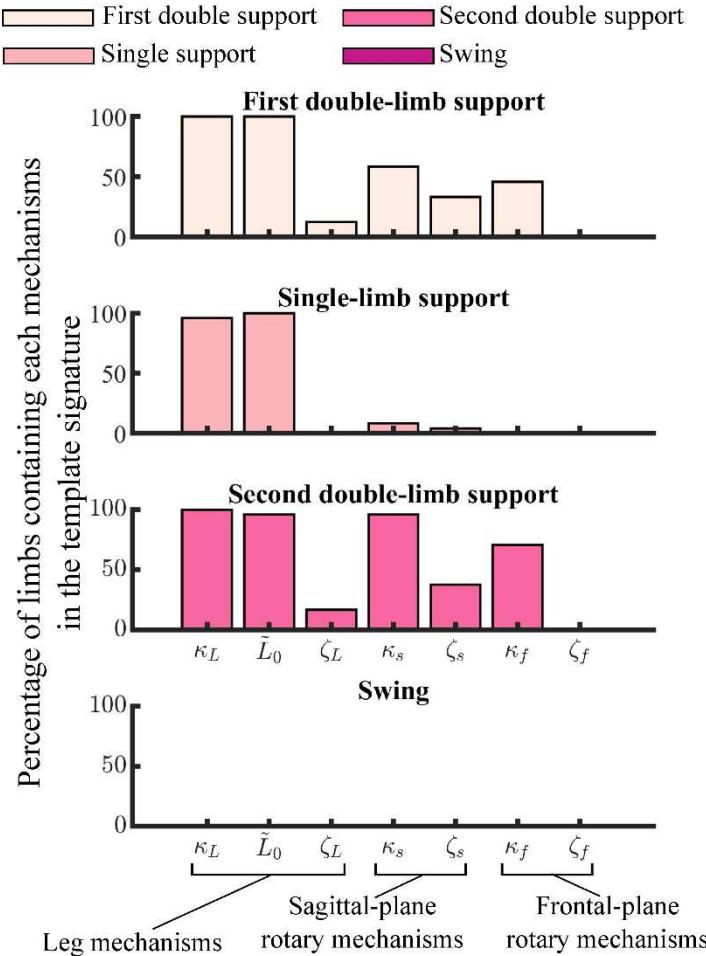


Figure 3 – The percentage of unimpaired limbs (24 limbs) whose template signatures contained each mechanism in each hybrid regime. Colors denote each hybrid regime. Mechanisms selected in a larger percentage of legs suggest common representations of COM dynamics, while less frequently-selected mechanisms reflect individual-specific features describing COM dynamics.

347

348 For each participant, template signatures were robust to stride-to-stride variations during single-limb
349 support: stiffness and leg length exhibited coefficients of variation (CVs) less than 0.021 ± 0.015 . In both

350 double-limb support phases, CVs of stiffness and sagittal-plane rotary damping ranged from 0.11 – 0.26.
351 Less frequently selected mechanisms in double-limb support – leg damping and frontal-plane damping –
352 were highly variable, with CVs ranging from 0.082 – 0.442.

353

354 Across unimpaired participants, single-limb support leg length was most consistent (CV = 0.03), while
355 leg stiffness was more variable (CV = 0.28; Figure 4; top). However, interindividual variability in double-
356 limb support leg and rotary stiffness was larger (CV = 0.43 – 1.94). Despite this variability, template
357 signatures reconstructed participants' COM accelerations well, 32-85% of the variance in 3D COM
358 accelerations (Figure 4; bottom).

359

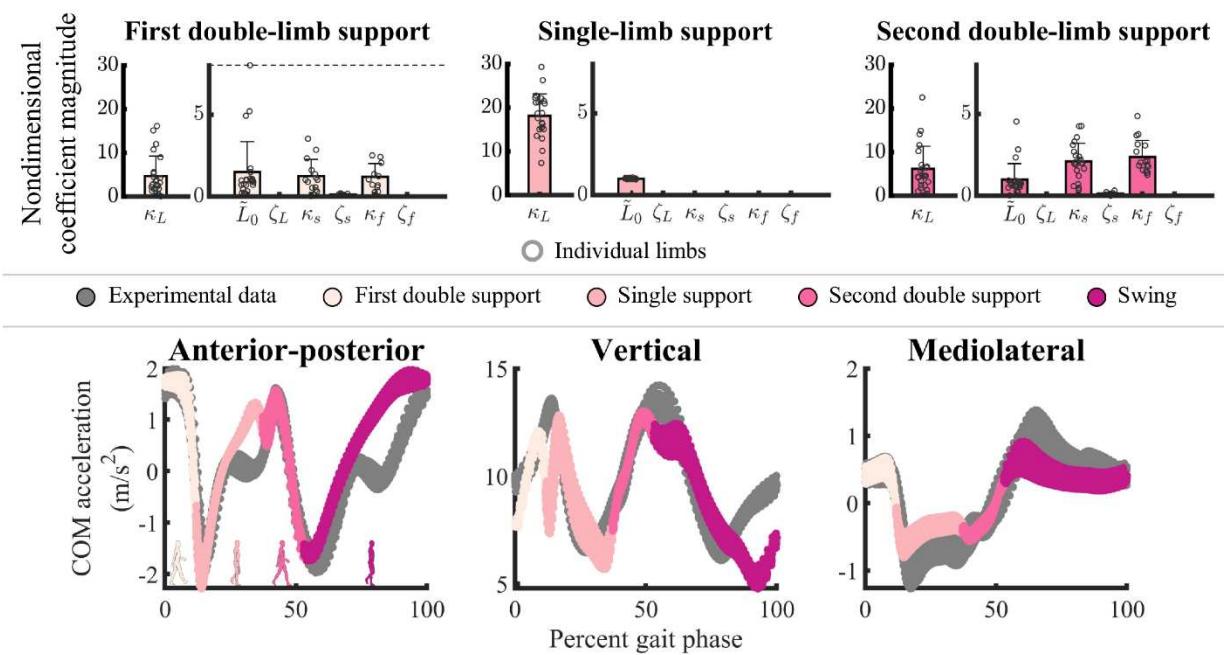


Figure 4 – Normalized template signatures (top) and reconstructed COM accelerations (bottom) for shoes-only walking in each hybrid regime. Top: Bars denote the average template signature ($\pm 1\text{SD}$) in single- and double-limb support. The small circles represent individual limbs. Note that we omitted mechanisms that were selected in less than 25% of participants. The dashed lines truncate large terms for clarity. Bottom: Experimental and predicted COM accelerations from the test dataset of an exemplary unimpaired participant. The gray dots denote the experimental accelerations, while the colors correspond to predictions in each hybrid regime.

360

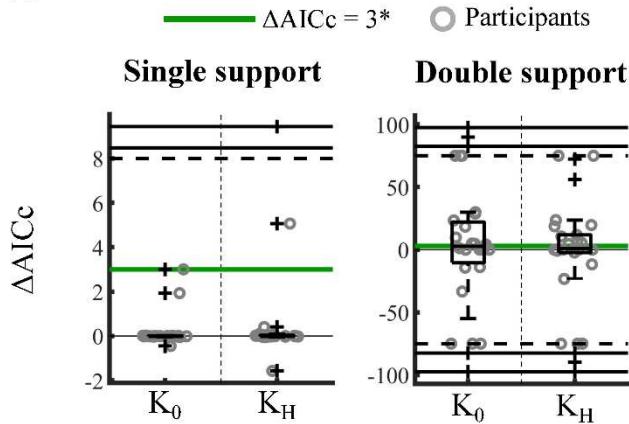
361 **Ankle exoskeleton impacts on unimpaired COM dynamics were small**

362 The shoes-only template signature structures reconstructed K_0 and K_H COM dynamics with similar
363 accuracy to the K_0 ($\Delta AICc = 0.2 \pm 0.7$; $p = 0.93$) and K_H ($\Delta AICc = 0.7 \pm 2.9$; $p = 0.24$) signature
364 structures, respectively, in single-limb support. Conversely, in double-limb support, shoes-only signatures
365 did not consistently reconstruct K_0 or K_H COM dynamics, with relative AICc scores ranging from $-118 \leq$
366 $\Delta AICc < 220$ (Figure 5; top). The negative relative AICc scores occurred when the shoes-only template
367 signature structure was never identified by Hybrid-SINDy in the exoskeleton conditions. Therefore, to
368 compare signature coefficients between exoskeleton conditions, we constrained each participant's
369 template signatures to that selected in the shoes-only condition.

370

371 Using this approach, we found that no unimpaired template signature coefficients changed significantly
372 when walking in shoes-only, zero-stiffness exoskeletons (K_0), and stiff exoskeletons (K_H) ($p > 0.05$).
373 Specific to our hypothesis, we found that neither ankle exoskeleton mass and frame (K_0) nor stiffness
374 (K_H) had a significant impact on leg stiffness in any gait phase ($p > 0.13$) (Figure 5; bottom).

(a)



$^* \Delta AICc \leq 3$ implies that shoes-only template signatures are plausible for walking with exoskeletons

(b)

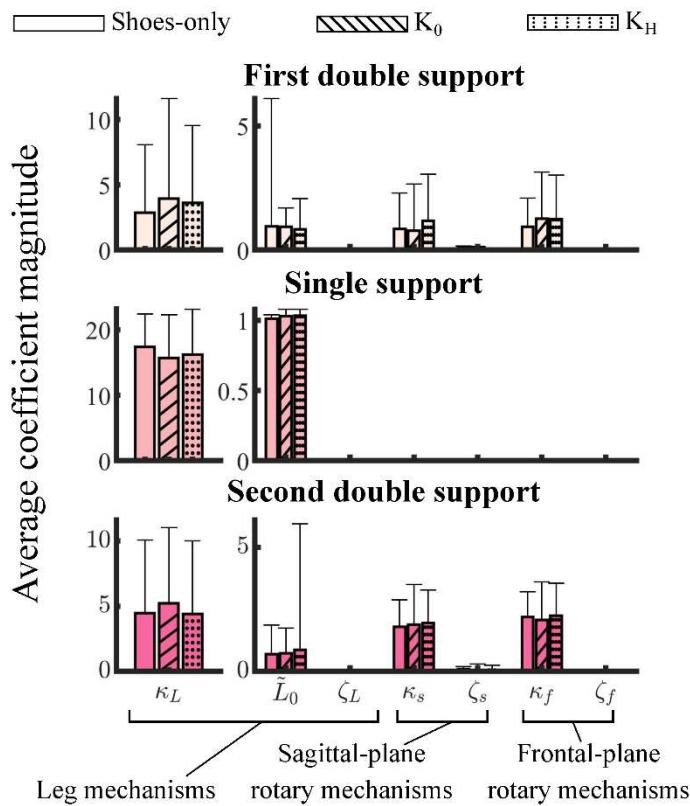


Figure 5 – (a) Relative AICc ($\Delta AICc$) between template signatures identified specifically for the K_0 and K_H conditions, and signatures constrained to the shoes-only signature structures. Positive $\Delta AICc$ values indicate that shoes-only signature structures were less plausible than those identified specifically for each condition. $\Delta AICc$ scores are shown for single-limb support (left) and double-limb support (right). $\Delta AICc \leq 3$ (green lines) denote plausible shoe template signature structures. For double-limb support,

large $\Delta AICc$ scores are truncated at $\Delta AICc = \pm 75$ for clarity. (b) Template signatures of walking in shoes only (solid bars), zero-stiffness (K_0 ; slashed bars) exoskeletons, and high-stiffness exoskeletons (K_H ; dotted bars) during single- and double-limb support. Bars represent the average (+1SD) template signature across participants. Leg stiffness (κ_L) is included on a separate subplot for clarity. The large variability in leg length during double-limb support is due one limb with a high resting length.

375

376 **Template signatures reveal altered COM dynamics post-stroke**

377 During shoes-only walking, the stroke survivor's template signature revealed distinct strategies for
378 accelerating the COM in double, but not single-limb support. The template signature structure was
379 symmetric between the paretic and non-paretic limbs during single-limb support, consisting of only leg
380 stiffness and resting length (Figure 6). However, the paretic limb was 31% stiffer than the non-paretic
381 limb. Conversely, template signature structure differed in double-limb support, with rotary stiffness
382 mechanisms improving descriptions of COM dynamics in only the paretic limb. During first double-limb
383 support, leg damping and rotary stiffness were selected in only the paretic limb. In second double-limb
384 support, both paretic and non-paretic rotary stiffness mechanisms were selected. Due to asymmetric
385 template signature structures, we did not compare signature coefficients in double-limb support.

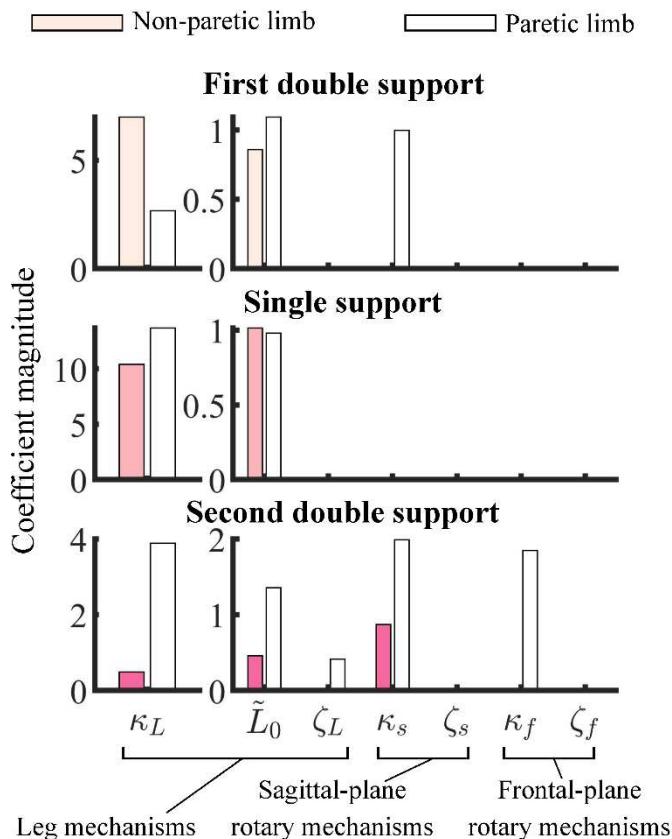


Figure 6 – Non-paretic (colored bars) and paretic (white bars) template signatures for one individual with post-stroke hemiparesis. This participant's template signatures differed in the paretic and non-paretic limbs in double-limb support, as shown by the zero-values for non-paretic sagittal-plane rotary stiffness in first double-limb support (top row) and the leg damping and frontal-plane stiffness in double-limb support (third row).

386

387 Ankle exoskeletons impacted post-stroke COM dynamics

388 The exoskeleton mass and frame (Shoe vs. K_0) primarily impacted the stroke survivor's paretic leg
389 stiffness and non-paretic leg rotary stiffness. Compared to walking in shoes-only, paretic leg stiffness was
390 11% greater in the zero-stiffness (K_0) exoskeleton condition in single-limb support and over twice as stiff
391 in double-limb support (Figure 7; slashed bars). Responses to the K_0 exoskeleton condition were
392 asymmetric, with the non-paretic leg being slightly (3%) less stiff in the K_0 condition than the shoes-only

393 condition in single-limb support, while sagittal-plane stiffness was 41% larger in second double-limb
394 support. Conversely, exoskeleton stiffness (K_0 vs. K_H) had small impacts on paretic-leg template
395 signatures. In the high-stiffness (K_H) exoskeleton condition, single-limb support paretic leg stiffness was
396 only 1% larger than in the K_0 condition (Figure 7; dotted bars). Both paretic and non-paretic leg rotary
397 stiffness were 11% greater in second double limb support than in the K_0 condition. Non-paretic leg
398 stiffness was 49% smaller than with zero-stiffness (K_0) exoskeletons in single-limb support.

399

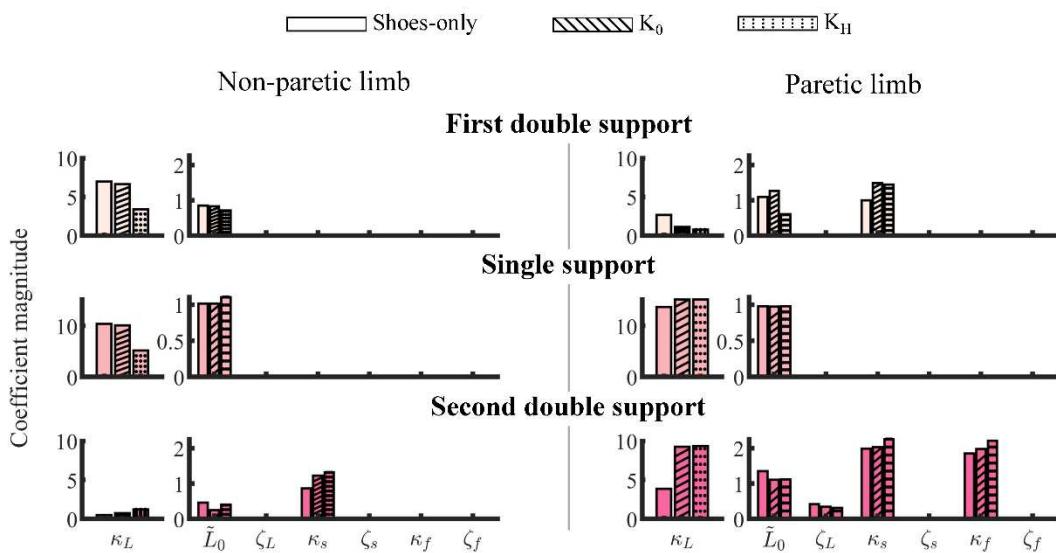


Figure 7 – Template signatures of a stroke survivor walking in the shoes-only (clear bars), zero-stiffness (K_0 ; slashed bars), and high-stiffness (K_H ; dotted bars) ankle exoskeleton conditions for the non-paretic (left) and paretic (right) limbs. Bars represent the average template signatures over 200 bootstrapped model fitting iterations. Colors correspond to each hybrid regime.

400

401 Discussion

402 Template signatures elucidate exoskeleton impacts on COM dynamics

403 We evaluated the impacts of passive ankle exoskeletons on subject-specific COM dynamics – described
404 by template signatures – using a recently-developed data-driven modeling framework, Hybrid-SINDy. To
405 our knowledge, this is the first study to quantitatively characterize the impacts of ankle exoskeletons on
406 COM dynamics. We used simulated SLIP to show that Hybrid-SINDy was robust to measurement noise
407 comparable to marker-based motion capture ($\eta = 1 \text{ mm}$). Therefore, template signature structure and
408 parameters identified by Hybrid-SINDy for human walking likely encode meaningful information about
409 COM dynamics. Contrary to our hypothesis, leg stiffness did not change in response to the ankle
410 exoskeletons in unimpaired adults. Further, template signatures identified from shoe walking were
411 plausible representations of COM dynamics when walking with ankle exoskeletons. These results suggest
412 that unimpaired strategies to accelerate the COM during walking are minimally impacted by passive ankle
413 exoskeletons, despite changes in kinematics and muscle activity [36]. Similarly, Collins and colleagues
414 (2015) observed small changes in total COM power with passive ankle exoskeletons compared to walking
415 in shoes-only, but larger changes in biological contributions to COM power [5]. However, the authors did
416 not explain how changes in COM energetics occurred. Our results suggest that these small changes in
417 COM power were accompanied by negligible changes in COM dynamics but were instead driven by
418 altered COM motion. However, maintaining COM dynamics with varying ankle exoskeleton properties
419 requires modulating joint or muscle-level (*i.e.*, execution-level) dynamics [5, 7, 36, 43]. Therefore,
420 quantifying subject-specific trade-offs between the modulation of task-level and execution level dynamics
421 may improve our understanding of responses to exoskeletons or other assistive devices.

422

423 Conversely, in one stroke survivor, ankle exoskeletons impacted COM dynamics. The addition of
424 exoskeleton mass and frame (K_0) increased paretic limb sagittal-plane rotary stiffness in first double-limb
425 support and leg stiffness in single-limb and second double-limb support. Increased paretic-leg stiffness is

426 likely due to the exoskeleton frame restricting ankle inversion, which the participant noted during data
427 collection. As the participant's paretic ankle was severely affected, restricting frontal-plane ankle motion
428 may have increased leg stiffness by stabilizing the ankle. This result emphasizes the importance of
429 quantifying the impacts of ankle exoskeleton properties beyond flexion stiffness, such as restriction to
430 frontal-plane ankle motion, in explaining exoskeleton impacts on gait following neurological injury [1, 5,
431 44].

432

433 The insensitivity of the paretic limb template signatures to exoskeleton stiffness during single-limb
434 support suggests that for this individual exoskeleton stiffness did not alter body weight support strategies
435 [45]. As the zero-stiffness exoskeletons increased leg stiffness, it is possible that resisting frontal-plane
436 ankle motion was the primary driver of changes in overall leg stiffness. That is, exoskeleton dorsiflexion
437 stiffness did not further increase the paretic ankle's contribution to body-weight support. Conversely,
438 during double-limb support, paretic-leg rotary stiffness increased compared to the zero-stiffness
439 exoskeleton condition. This change in COM dynamics suggests that exoskeletons primarily altered
440 dynamics impacting forcing transverse to the leg, which primarily contribute to propulsion and
441 stabilization [46]. Unlike single-limb support, exoskeleton stiffness in terminal stance may enable
442 increased propulsive force in the paretic limb, potentially explaining rotary mechanisms' increased
443 contributions to gait. Further, the ankle plantarflexors accelerate the COM laterally, such that increased
444 paretic plantarflexion torque capacity may impact frontal-plane dynamics as well [47, 48]. Note that,
445 while these results indicate that our participant's task-level dynamics corresponding to stabilization and
446 propulsion were impacted by exoskeleton stiffness, this result is likely to differ between stroke survivors
447 [2]. Specifically, exoskeleton impacts on COM dynamics may depend on an individual's ability to
448 convert exoskeleton assistance into functionally relevant forcing. While limited to a case study, our
449 results highlight the potential contributions of ankle exoskeleton frame and stiffness on post-stroke COM
450 dynamics that can be described by template signatures.

451

452 **Template signatures quantify inter-individual and inter-limb differences in COM**

453 **dynamics**

454 Both unimpaired and post-stroke template signatures highlighted inter-individual and inter-limb
455 differences in COM dynamics. While SLIP mechanisms described COM dynamics in all unimpaired
456 limbs, rotary mechanisms were selected in only 50% of limbs during double-limb support. The uniform
457 selection of SLIP mechanisms from data is consistent with common template walking models and
458 supports the common perspective that elastic legs are foundational mechanisms for describing COM
459 accelerations during walking across individuals [13, 21, 22, 26, 49]. Our observation that rotary
460 mechanisms were not critical to reconstructing COM accelerations for all individuals may explain their
461 less frequent application in template walking models [14, 20]. Note that each template signature
462 mechanism describes characteristic coordination patterns between leg kinematics and COM accelerations.
463 It is possible SLIP mechanisms describe coordination patterns necessary for stable or efficient walking,
464 while rotary mechanisms describe coordination patterns that have smaller, more individualized impacts
465 on gait. Determining which individual-specific characteristic patterns are predictive of exoskeleton
466 responses or walking function represents important areas of future research.

467

468 Template signatures also revealed compensation strategies in one stroke survivor. Consistent with
469 template-based studies in children with cerebral palsy, the stroke participant's paretic leg was stiffer than
470 the nonparetic leg in single- and second double-limb support [29, 30]. Increased paretic leg stiffness may
471 reflect a more pendular gait post-stroke and increased paretic-leg co-contraction [50]. In first double-limb
472 support, sagittal-plane rotary stiffness and damping suggest altered braking dynamics during loading
473 response. This possibility is consistent with experimental observations that the paretic-leg does greater
474 negative work than the non-paretic limb during loading response in individuals post-stroke [12]. These
475 results highlight the functional interpretability of data-driven template signatures and the importance of

476 personalized parameterizations of COM dynamics for elucidating inter-limb differences in COM
477 dynamics.

478

479 **Innovation of Hybrid-SINDy for quantifying human gait**

480 Our use of the Hybrid-SINDy algorithm ([33, 34]) was essential to elucidating individual-specific and
481 limb-specific changes in COM dynamics with ankle exoskeletons. By automatically identifying and
482 selecting different first principles-based representations of COM dynamics, we rapidly compared
483 competing hypotheses about human gait dynamics. Conversely, manually comparing all possible template
484 signatures for our fourteen-dimensional mechanism library would require a combinatorially-large number
485 of models to be generated, fit, and compared. Using a principled approach to model identification and
486 selection, Hybrid-SINDy selected dynamics that are consistent with literature in a fraction of the time
487 required to manually compare each candidate model.

488

489 To our knowledge, only one other study in bipedal locomotion has taken a similar approach [51]. The
490 authors performed a multi-layer optimization to select human-like template dynamics for walking robot
491 controllers. Our approach extends their work by using information criteria to enable multiple highly-
492 plausible representations of walking dynamics to be selected and analyzed for an individual [35, 40]. For
493 most participants, the Hybrid-SINDy algorithm selected a unique template signature, indicating that
494 alternative hypothetical representations of COM dynamics lacked strong statistical support. By
495 identifying multiple plausible template signatures for an individual, future experiments may be designed
496 to better differentiate between plausible signatures. To our knowledge, this work is the first to apply
497 Hybrid-SINDy to real data and the first to apply it to human gait.

498

499 Since reduced-order models are often sensitive to measurement noise and inevitably incorrect model
500 dynamics, we used a simulated SLIP with known dynamics to evaluate the extent to which Hybrid-

501 SINDy correctly identified template signatures during walking. Hybrid-SINDy performed as expected in
502 the noise-free condition, accurately identifying single-limb support and swing dynamics, except for small
503 damping terms. Terms with small contributions to system behavior, such as damping in this study, may be
504 omitted by the AICc in favor of parsimony [35, 40]. An important implication of this result is that our
505 approach will not always select the true system dynamics but will select terms critical to describing
506 salient behavior of the system. Therefore, designing experimental protocols capable of probing candidate
507 mechanisms is critical to accurately identifying walking dynamics from data. For example, gait patterns
508 that induced large leg velocities may result in damping terms having a larger impact on COM dynamics
509 and being correctly selected by Hybrid-SINDy. Additionally, reduced simulated SLIP template signature
510 accuracy in double-limb support and with increasing noise magnitude highlights the importance of
511 collecting long time-series datasets to ensure that clusters are small enough to span only a single hybrid
512 regime but large enough to remain robust to sensor noise [33].

513

514 **Challenges and limitations**

515 This study's limitations motivate future developments for data-driven modeling of human locomotion.
516 First, human gait dynamics are continuously phase-varying rather than hybrid, such that template
517 signature parameters varied continuously across clusters. However, preliminary analyses found candidate
518 models' abilities to reconstruct COM dynamics diverged near the transitions between single and double-
519 limb support, suggesting that changes in dynamics between foot contact configurations are larger than
520 changes within configurations. Consequently, pre-defining hybrid regimes based on contact configuration
521 was both appropriate and consistent with existing hybrid template models of walking [20, 21, 26, 33].
522 Second, we evaluated only a subset of mechanisms describing COM dynamics, which may not encode all
523 important features of human gait. However, we chose a broad class of physically-meaningful mechanisms
524 from literature ([18, 20-22, 26, 28]) that should motivate future studies using larger mechanism libraries
525 describing COM dynamics. For example, our approach may be extended to include elements describing

526 assistive devices or joint-level mechanisms. Finally, our limited sample size may have masked
527 exoskeleton impacts on unimpaired gait. Particularly, rotary stiffness appeared to increase slightly with
528 exoskeletons but did not reach significance. Similarly, our post-stroke case study only provides a proof of
529 concept highlighting the potential of ankle exoskeletons to elicit changes in COM dynamics. Evaluating
530 template signatures on larger unimpaired and post-stroke cohorts may provide additional insight into how
531 COM dynamics change with ankle exoskeletons.

532

533 [Extensions and applications](#)

534 Integrating principles from biomechanics and neuromechanics with state-of-the-art *interpretable* data-
535 driven modeling techniques may accelerate discovery in mechanisms underlying diverse movement
536 patterns in human locomotion. Broadly, these methods may be applied to any intervention or walking
537 condition and may inform the design of prosthetic limbs and other assistive devices. A natural extension
538 of this work is to investigate the impacts of powered exoskeletons on COM dynamics. Changes in gait
539 kinematics and kinetics with our passive ankle exoskeletons were small compared to those elicited by
540 powered exoskeletons, which may elicit larger changes in COM dynamics [2, 6, 36]. As powered
541 exoskeleton research shifts its focus from reducing metabolic demand to assisting other locomotion
542 objectives, such as stability or agility, understanding changes in COM dynamics with exoskeletons may
543 provide insight into how individuals leverage exoskeletons to realize these goals [52]. Particularly in a
544 clinical context, quantifying changes in COM dynamics with exoskeletons may improve our
545 understanding of the diverse responses to exoskeletons observed in individuals with neurological injuries
546 [1-3]. For example, McCain and colleagues (2019) identified limb orientation as a potential mechanism
547 impacting the conversion of exoskeleton torque to forward propulsion post-stroke [2]. Our results in one
548 stroke survivor suggest that characterizing altered coordination between leg orientation and ground
549 reaction forces (*i.e.*, template signatures) may unveil sub-classes of exoskeleton responses, enabling
550 device customization to assist individuals in each sub-class. Finally, using similar physics-informed data-

551 driven approaches to *anchoring* template signatures in physiological detail may reveal biological
552 mechanisms that characterize responses to exoskeletons or other assistive devices [14, 16].

553

554 **Conclusions**

555 We quantified changes in subject-specific COM dynamics in response to passive ankle exoskeletons
556 using a novel physics-informed data-driven modeling framework, Hybrid-SINDy. Hybrid-SINDy
557 successfully constructed characteristic template signatures during walking by identifying salient
558 mechanisms describing an individual's COM accelerations. Unimpaired adults maintained nearly
559 invariant COM dynamics with ankle exoskeletons, while exoskeletons altered COM dynamics in an
560 individual post-stroke. These findings provide insight into task-level control of walking dynamics with
561 ankle exoskeletons and extend a powerful data-driven modeling tool to human locomotion.

562

563

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