

1 **Validity and sensitivity of an inertial measurement unit-driven biomechanical model of**
2 **motor variability for gait**

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Abstract

Motor variability in gait is frequently linked to fall risk, yet field-based biomechanical joint evaluations are scarce. We evaluated the validity and sensitivity of an inertial measurement unit (IMU)-driven biomechanical model of joint angle variability for gait. Fourteen healthy young adults completed seven-minute trials of treadmill gait at several speeds and arm swing amplitudes. Joint kinematics were estimated by IMU- and optoelectronic-based models using *OpenSim*. We calculated range of motion (ROM), magnitude of variability (meanSD), local dynamic stability (λ_{\max}), persistence of ROM fluctuations (DFA α), and regularity (SaEn) of each angle over 200 continuous strides, and evaluated model accuracy (e.g., RMSD: root mean square difference), consistency (ICC_{2,1}: intraclass correlation), biases, limits of agreement, and sensitivity to within-participant gait responses (effects of Speed and Swing). RMSDs of joint angles were 1.7–7.5° (pooled mean of 4.8°), excluding ankle inversion. ICCs were mostly good–excellent in the primary plane of motion for ROM and in all planes for meanSD and λ_{\max} , but were poor–moderate for DFA α and SaEn. Modeled Speed and Swing responses for ROM, meanSD, and λ_{\max} were similar. Results suggest that the IMU-driven model is valid and sensitive for field-based assessments of joint angles and several motor variability features.

Keywords: gait, human movement, inertial measurement unit, joint kinematics, local dynamic stability, musculoskeletal model, OpenSim, persistence, regularity, stride-to-stride variability

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1. Introduction

21 Motor variability, the natural variability in sensorimotor actions [1], is well-linked to
22 walking-related fall risk. Elderly fallers exhibit greater stride-to-stride variability in
23 spatiotemporal outputs (e.g., stride time) compared to non-fallers [2,3]. This difference may
24 emerge from altered stride-to-stride joint angle patterns that have been observed with older age,
25 including lower local dynamic stability [4], lower regularity [5], and a shift in the magnitude of
26 variability in ankle motion from the sagittal to the frontal plane [6]. Monitoring the variability of
27 joint angles in aging adults at a larger scale could help better understand the salient elements of
28 stride-to-stride control that predict falls, which could better individualize fall-prevention
29 interventions. However, measurements of joint angles (and variability from stride to stride)
30 typically rely on optoelectronic motion capture systems that are expensive, require training and
31 expertise to operate, and involve intensive data acquisition and processing procedures.
32 Optoelectronic motion capture of gait is also restricted to treadmills [4,5] or short distances
33 overground [6], which may not fully replicate the stride-to-stride variability that occurs over
34 longer distances and durations overground. Using this motion capture approach, large-scale
35 evaluations of motor variability in realistic and clinically relevant gait scenarios are infeasible.

36 Wearable inertial measurement units (IMUs) offer an alternative technology for
37 estimating joint angles that can address these limitations of optoelectronic motion capture. IMUs
38 have been used to estimate joint kinematics since 1990 [7], with recent work showing that IMU-
39 based kinematic models of lower-limb activities achieve absolute differences (i.e., accuracies)
40 ranging from 1 to 10° relative to optoelectronic-based models [8–14] and good–excellent
41 consistencies in the sagittal plane timeseries [8,11,15,16]. Some IMU-based kinematic models
42 are built based on machine learning approaches [12], but most others typically involve (i)

43 estimating the IMU orientation by fusing accelerometer and gyroscope data, (ii) estimating the
44 anatomical segment orientation by applying a sensor-to-segment calibration, and (iii) calculating
45 joint angles as the relative orientation between reference frames fixed to adjacent body segments
46 (see [17] for review). A particular obstacle to the longer-duration recordings necessary to
47 measure stride-to-stride kinematics is preventing IMU-sensor orientation drift attributed to
48 sensor fusion. Strap-down integration of an IMU on a non-stabilized segment amplifies random
49 noise in linear accelerations and angular velocities, leading to drift in the estimated orientations.
50 Many sensor fusion algorithms address horizontal drift by incorporating data on the Earth's
51 magnetic field detected by the magnetometer [18–22]; however, detection of the magnetic field
52 is disturbed locally by ferromagnetic materials [23]. Beyond the magnetometer, drift corrections
53 are possible over short durations by adding zero-velocity updates [24] and over short distances
54 by adding localization using technologies like ultrawideband [10], but these approaches do not
55 provide a solution for long durations or distances.

56 Anatomical joint constraints in the underlying kinematic model can help to mitigate drift
57 over long durations and distances. Using the *OpenSense* toolkit to compute inverse kinematics of
58 a biomechanical model with IMU inputs, Al Borno et al. [14] recently demonstrated root mean
59 squared differences (RMSD) of 3–6° for IMU-based hip flexion, hip abduction, knee flexion,
60 and ankle dorsiflexion angles relative to optoelectronic motion capture, with near-zero drift
61 (from 0.14 to 0.17°/min) over ten minutes of walking. This supports the earlier findings of Kok
62 et al. [25] and provides a new open-source option for constructing an IMU-based kinematic
63 model. This inverse kinematics approach to solving joint angles has the added benefit of
64 mitigating experimental (i.e., non-biological) noise [26], which could particularly benefit
65 evaluations of motor variability. Although constrained optimization problems like inverse

66 kinematics can require high computation time to solve, recent work from Slade et al. [13]
67 demonstrated that a real-time IMU-based solution is possible. They reported RMSD in joint
68 angles of approximately 5°, a difference accepted as reasonable for many clinical applications
69 [27]. The *OpenSense* extension of *OpenSim* [28,29] provides the first open-source platform for
70 IMU-based biomechanical modelling and a free alternative to cost-prohibitive and closed-source
71 commercial models [18]. Because Al Borno et al. [14] used the magnetometer to calculate drift-
72 free kinematics and Slade et al. [13] reported kinematic drifts in their magnetometer-free solution,
73 it remains unclear whether a magnetometer-free, IMU-based biomechanical model can provide
74 accurate joint angles during gait beyond one minute duration. Furthermore, it is unknown
75 whether IMU-based biomechanical models are valid for evaluating stride-to-stride variability or
76 are sufficiently sensitive to changes in gait kinematics for evaluating fall risk.

77 The goal of this study was to validate a magnetometer-free, open-source, IMU-based
78 biomechanical model of joint angles and stride-to-stride variability for a moderate duration of
79 continuous gait. We determined the concurrent validity of IMU-based and optoelectronic-based
80 model measurements of joint angles and joint angle variability from the trunk down using
81 *OpenSim*, and determined the sensitivity of discrete measurements of joint angles and joint angle
82 variability to different gait conditions. We have made the data underlying the findings of this
83 study freely available online, enabling others to reproduce and extend our work.

84 **2. Methods**

85 **2.1 Participants**

86 Fourteen healthy young adults (7 males, 7 females) were recruited as a convenience
87 sample from the Ottawa, Canada area. This sample size (n = 12, with 2 extra to account for

88 possible data attrition) was determined *a priori* using G*Power [30] based on our sensitivity
89 analyses and reflects the number of participants needed to detect a large effect size (partial $\eta^2 \geq$
90 0.14) for a within-group factor with three measurement levels (Speed: preferred, slow, fast;
91 Swing: preferred, active, bound) at a power of 0.80 and an α of 0.05. Participants were excluded
92 if they had a musculoskeletal injury in the preceding 6 months, or any chronic neurological or
93 orthopaedic disorders. Participants all provided written informed consent to the study, which
94 followed the Declaration of Helsinki and was approved by the University of Ottawa Research
95 Ethics Board (H-01-21-6261).

96 **2.2 Instrumentation**

97 Following informed consent, the participants donned spandex motion capture pants and
98 their own athletic shoes. Participants were then instrumented for optoelectronic- and IMU-based
99 motion capture (Figure 1). An 11-camera optoelectronic system (Vantage, Vicon, Oxford, UK)
100 sampled trajectories of spherical retroreflective markers at 120 Hz using Nexus 2.11 (Vicon,
101 Oxford, UK). Markers were placed on the participant's body using double-sided tape, following
102 the marker locations used with the full-body model for gait simulations in *OpenSim* from
103 Rajagopal et al. (Link: https://simtk.org/projects/full_body) [31]. Anatomical markers were
104 placed on each wrist (radial and ulnar styloid process), on each elbow (medial and lateral
105 epicondyle), on the trunk (left and right acromion process, right clavicular head, spinous process
106 of C7), on the pelvis (left and right anterior superior iliac spine, left and right posterior superior
107 iliac spine), on each knee (medial and lateral femoral condyle), on each ankle (medial and lateral
108 malleolus), and on each foot (heel, 1st and 5th metatarsal head). The dynamic marker set was
109 modified to add redundancy such that rigid-body clusters of four markers (rather than three) were
110 positioned using Velcro straps on the trunk and each forearm, arm, thigh, and shank. Anatomical

111 markers on the feet, pelvis, and the lateral malleolus markers on the shank doubled as dynamic
112 markers for these segments.

113 IMU-based motion capture was performed using a platform of eight wearable sensors
114 (Dot, Xsens, Enschede, Netherlands) and a mobile application (Xsens Dot Precision Motion
115 Tracking) for synchronized acquisition of the raw accelerometer and gyroscope data from each
116 sensor. IMU sensors were positioned on the feet (top of the shoe), shanks (anterior aspect, distal
117 and immediately above the malleoli), thighs (anterior aspect, around the largest circumference),
118 pelvis (posterior aspect under the posterior superior iliac spines), and trunk (posterior aspect at
119 the level of the sternum). IMUs were oriented such that the positive xyz axes in the sensor frame
120 in anatomical position were directed leftward, upward, and forward, respectively. Raw
121 accelerations and angular velocities were sampled at 60 Hz using the mobile application in data
122 logging mode.



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Figure 1. Participant in standing pose on the treadmill showing the dynamic marker set, inertial measurement unit positions (strapped to segments, circled in orange), and the wooden block.

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2.3 Experimental procedure

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Following instrumentation, preferred gait speed was identified according to the procedure of Dingwell and Marin [32]. With the participant blinded to the speed and walking slowly on the treadmill, gait speed was progressively increased until they reported that the speed was “faster than preferred”. Speed was then progressively decreased until they reported that the speed was “slower than preferred”. This sequence was then repeated three times, with the average of the six speeds defined as the preferred gait speed.

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Following a static calibration of the anatomical markers with the participant in standing pose, the participant completed five gait trials on the treadmill (Horizon Fitness, WI, USA). Each trial began with a 30-second procedure to warm up the sensors, with the feet oriented on the

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136 treadmill using a wooden block to minimize inter-trial and inter-individual differences in foot
137 excursion posture (Figure 1). The participant stood quietly for the first 5 seconds, leaned forward
138 for the next 10 seconds, then returned to quiet standing for the final 15 seconds. After this
139 baseline procedure, the block was removed and the participant completed seven minutes of
140 walking. This sequence was repeated for five trials, each under a different gait condition that
141 varied by gait speed and/or arm swing magnitude: 1) preferred gait speed and arm swing; 2) 70%
142 preferred gait speed, preferred arm swing; 3) 130% preferred gait speed, preferred arm swing; 4)
143 preferred gait speed, active arm swing (the participant was instructed to swing their arms such
144 that forward swing peaked when the arm was horizontal); and 5) preferred gait speed, arms
145 bound to the torso (using straps across the arms and across the elbows). Different gait speeds and
146 arm swing amplitudes were evaluated since they have been shown to alter stride-to-stride
147 variability patterns in gait [33–37], providing a basis for exploring the sensitivity of the IMU-
148 based model. Condition order was randomized, the participant rested for a minimum of three
149 minutes between trials, and optoelectronic and IMU data were continuously sampled during each
150 trial.

151 **2.4 Data Analysis**

152 **2.4.1 Optoelectronic-based biomechanical modelling.** Marker trajectories were labeled,
153 gap-filled with a Woltring spline [38], and low-pass filtered at 10 Hz using Nexus (v2.11, Vicon
154 Inc., Oxford, UK). Filtered trajectories were then imported into *OpenSim* v4.1 [29] and used to
155 simulate motion of a full-body model containing 37 degrees of freedom (DOF) and 80 muscle–
156 tendon units actuating the lower limbs [31]. This model includes a 3-DOF trunk (relative to the
157 pelvis), a 6-DOF pelvis (relative to ground), 3-DOF hips, 1-DOF knees, 2-DOF ankles, and 1-
158 DOF toes. The 1-DOF toe joints were locked since toe motion was not recorded by IMUs. The

159 model was scaled to the participant using the positions of the anatomical markers in the static
160 trial; joint angles in each trial were then computed via an inverse kinematic analysis that
161 minimized the least-squared distance between each pair of experimental and model markers.
162 Upper-limb markers were not included in the analysis since no IMUs were placed on the upper
163 limbs and applying weights to the upper-limb markers that equaled weights of lower-limb
164 markers did not influence the inverse kinematics solution (Supplementary material: Figure S1).
165 Marker weights were manually selected to minimize the root-mean-square error over all marker
166 pairs, resulting in equal weights except for weights of twice the magnitude for markers on the
167 acromion processes (trunk), anterior and posterior superior iliac spines (pelvis), and lateral
168 malleoli (shanks).

169 **2.4.2 IMU-based biomechanical modelling.** Using Matlab (R2020b, The MathWorks
170 Inc., MA, USA), raw linear accelerations and angular velocities were fused offline to calculate
171 sensor orientations using the magnetometer-free algorithm of Madgwick et al. [19]. Orientation
172 drifts were then removed using a detrending procedure. Beginning at the 20-second timestamp (5
173 seconds after the participant had completed the forward lean and was standing quietly) and
174 ending at trial completion, quaternions were converted to ZYX Euler angles and fit to a function
175 using ‘polyfit’. Linear drift was identified as slope less than -0.0010 rad/s or greater than 0.0010
176 rad/s and removed from the Euler angle signal using ‘detrend’, then Euler angles were converted
177 back to quaternion representation. An example showing the orientation of a sensor before and
178 after drift removal is provided in the supplementary material (Figure S2).

179 Detrended sensor quaternions were imported into *OpenSim* using the *OpenSense* toolkit
180 to simulate motion of the same full-body biomechanical model [31]. Sensor-to-segment
181 registration was performed to associate the orientation of each sensor with the corresponding

182 segment in the model; specifically, each thigh, shank, and foot sensor was registered,
183 respectively, to each femur, tibia, and talus body segment. Sensor orientations were converted
184 from their local coordinate systems to the *OpenSim* coordinate system using the following
185 sequence of body-fixed rotations: 180° about x, then 90° about y, and finally -90° about z. IMU
186 segment frames were identified based on the standing pose at the start of each gait trial: fixed
187 rotational offsets were applied to recorded IMU sensor frames based on the segment frames of
188 the biomechanical model in a neutral standing pose (i.e., joint flexion of 0°), with heading offsets
189 applied to individual IMU sensor frames to match the average heading and align with the
190 anterior–posterior axis of the biomechanical model [13,14]. As with the optoelectronic-based
191 model, joint angles in each trial were calculated via inverse kinematics; for the IMU-based
192 model, the solver minimized axis-angle differences between the IMU segment orientations and
193 IMU sensor orientations [14]. We compensated for differences between the initial pose of the
194 optoelectronic and IMU models by offsetting optoelectronic-based joint angle timeseries by a
195 constant to match the neutral standing pose of the IMU model. In contrast with the single ankle
196 dorsiflexion/plantarflexion DOF modeled previously [13,14], we chose to also model ankle
197 inversion/eversion since both sagittal- and frontal-plane ankle compensations are relevant to gait
198 of aging adults [6]. We explored optoelectronic- and IMU-based inverse kinematic solutions of
199 the 1-DOF and 2-DOF ankle models and confirmed that the additional ankle DOF did not affect
200 the inverse kinematics solutions for other lower-limb joint angles (Supplementary material:
201 Figure S1). Example animations of the optoelectronic- and IMU-based biomechanical models
202 can be viewed in the supplementary material (Videos S1–S4).

203 **2.5 Calculation of kinematic outcomes.** Timeseries of optoelectronic- and IMU-modeled joint
204 angles and linear velocities of the foot segments were exported to Matlab. Strides were

205 partitioned by the first and subsequent heel strike events, identified from the tri-dimensional
206 linear velocity vectors of the calcaneus segments. Linear velocities were low-pass Butterworth
207 filtered with zero lag (10 Hz cutoff, fourth-order) and the Euclidean norm linear velocity vector
208 was then calculated. From the Euclidean norm linear velocity, heel strike events were identified
209 as the local minima immediately following each local maximum (Supplementary material:
210 Figure S3). After removing strides in the first 30 seconds to ensure gait was at steady state, the
211 raw and time-normalized series (101 points per stride: 0–100%) of joint angles were analyzed for
212 the subsequent 200 strides. (This was the minimum number of synchronized strides available for
213 all trials and participants.) Five outcomes were then calculated for each joint angle, as described
214 below.

215 **2.5.1 Range of motion (ROM).** The difference between the maximum and minimum
216 angles for each stride in the time-normalized series. The mean value across strides was computed.

217 **2.5.2 Mean standard deviation (meanSD).** A measure of the absolute magnitude of
218 variability, meanSD was calculated using the time-normalized series. SD was calculated across
219 all strides for each time point ($n = 101$ points) and the mean SD across time points (meanSD)
220 was computed.

221 **2.5.3 Maximum finite-time Lyapunov exponent (λ_{\max}).** A measure of local dynamic
222 stability, λ_{\max} (i.e., the local divergence exponent) was calculated using the continuous series.
223 The continuous series was normalized to 20,000 points (100 per stride on average), then λ_{\max} was
224 computed with 5 embedding dimensions at a lag of 10 points from 0–0.5 strides (50 points)
225 [4,39]. λ_{\max} measures the local divergence of neighbouring trajectories with higher positive
226 values indicative of higher divergence and lower local dynamic stability.

227 **2.5.4 Detrended fluctuation analysis scaling exponent (DFA α).** A measure of
228 statistical persistence, DFA α was calculated as the fluctuation in ROM across strides, computed
229 as previously described [40,41] and quantifying the extent to which ROM fluctuations
230 statistically persist. DFA α is non-negative and unitless, with values greater than 0.5 indicating
231 persistence (i.e., a fluctuation is typically followed by a fluctuation in the same direction), values
232 less than 0.5 indicating anti-persistence (i.e., a fluctuation is typically followed by a fluctuation
233 in the opposite direction), and values around 0.5 indicating no correlation between consecutive
234 fluctuations [42].

235 **2.5.5 Sample entropy (SaEn).** A measure of regularity, SaEn was calculated using the
236 continuous series, 2 embedding dimensions, and a 0.15 tolerance distance [43]. SaEn can be
237 investigated at several scales using a multiscale function; we selected a scale factor of 4 which is
238 believed to be the approximate value where entropy of physiological signals stabilizes during
239 self-selected slow, normal (usual), and fast walking speeds [43]. To compensate for the influence
240 of sampling frequency [44], IMU-based joint angles were resampled at 120 Hz to match the
241 optoelectronic system. SaEn is non-negative and unitless, with higher values indicating lower
242 regularity.

243 **2.6 Statistical Analyses**

244 **2.6.1 Analyses of IMU-model validity.** For each degree of freedom, we assessed
245 concurrent validity of the IMU-based joint angles and angle outputs relative to the
246 optoelectronic-based joint angles and angle outputs for 1000 strides (5 gait conditions \times 200
247 strides). Mean RMSD was calculated for the time-normalized joint angle series as well as for
248 each outcome variable. For the timeseries analysis, relative difference was calculated as the

249 coefficient of variation of RMSD relative to the optoelectronic-based ROM (CV_{rom}); for the
250 outcome variable analysis, relative difference was calculated relative to the optoelectronic-based
251 mean (CV_{mean}). Intraclass correlation coefficients ($ICC_{2,1}$) and Bland-Altman plot metrics (IMU–
252 optoelectronic measurement bias, 95% limits of agreement) were computed to examine
253 consistency and agreement between outcomes. $ICC_{2,1}$ values less than 0.40, from 0.40 to 0.59,
254 from 0.60 to 0.74, and greater than or equal to 0.75 were interpreted as poor, fair, good, and
255 excellent consistency, respectively [45]. Biases indicated whether IMU-based outcomes were
256 overestimated (positive bias) or underestimated (negative bias) on average.

257 **2.6.2 Analyses of IMU-model sensitivity.** For each degree of freedom and outcome, we
258 assessed the sensitivity of the IMU model to detect the same within-participant changes as the
259 marker-based model by conducting repeated measures ANOVAs on each model to test for
260 effects of gait speed (Speed: preferred, 70% preferred, 130% preferred) and arm swing (Swing:
261 preferred, active, bound). Greenhouse-Geisser corrections were applied if sphericity was violated,
262 and critical alpha was set to 0.027 using the Benjamini-Hochberg procedure to account for false
263 discovery rate due to multiple comparisons [46] (240 p-values: 2 models \times 2 statistical effects \times
264 12 angles \times 5 outcomes). Post-hoc tests, comparing each condition to preferred speed and
265 preferred arm swing gait, were made with Bonferroni corrections ($p < 0.05$).

266 **3. Results**

267 **3.1 Participant characteristics**

268 Participant height, mass, and BMI (mean \pm SD) averaged 1.72 ± 0.07 m, 69.6 ± 14.2 kg,
269 and 23.5 ± 3.9 kg/m², respectively. Gait at preferred speed averaged 1.12 ± 0.18 m/s (range:

270 0.72–1.50), at 70% preferred speed averaged 0.79 ± 0.13 m/s (range: 0.50–1.05 m/s), and at
271 130% preferred speed averaged 1.46 ± 0.24 m/s (range: 0.93–1.95).

272 **3.2 IMU-model validity**

273 **3.2.1 Timeseries.** Mean values are presented in Table 1, with ensemble averaged curves
274 for the preferred speed condition displayed in Figure 2. For consistency, angles in the sagittal,
275 frontal, and transverse planes will be described as flexion/extension (FE), abduction/adduction
276 (AA), and internal/external rotation (IE), respectively. Based on values pooled across conditions,
277 mean RMSD was less than 5° for all trunk angles, pelvis FE, hip FE, and knee FE, with all other
278 angles except ankle AA approaching the 5° threshold. RMSD pooled across conditions and
279 angles was 5.3° , which dropped to 4.8° when ankle AA was excluded. RMSDs were consistent
280 across 200 consecutive strides, showing that IMU-modeled joint angles did not drift
281 (Supplementary material: Figure S4). CV_{rom} averaged 25.4% across angles, being lowest in the
282 transverse plane for the trunk (16.2%) and pelvis (12.8%), and lowest in the sagittal plane for the
283 hip (9.2%), knee (6.4%), and ankle (17.1%).

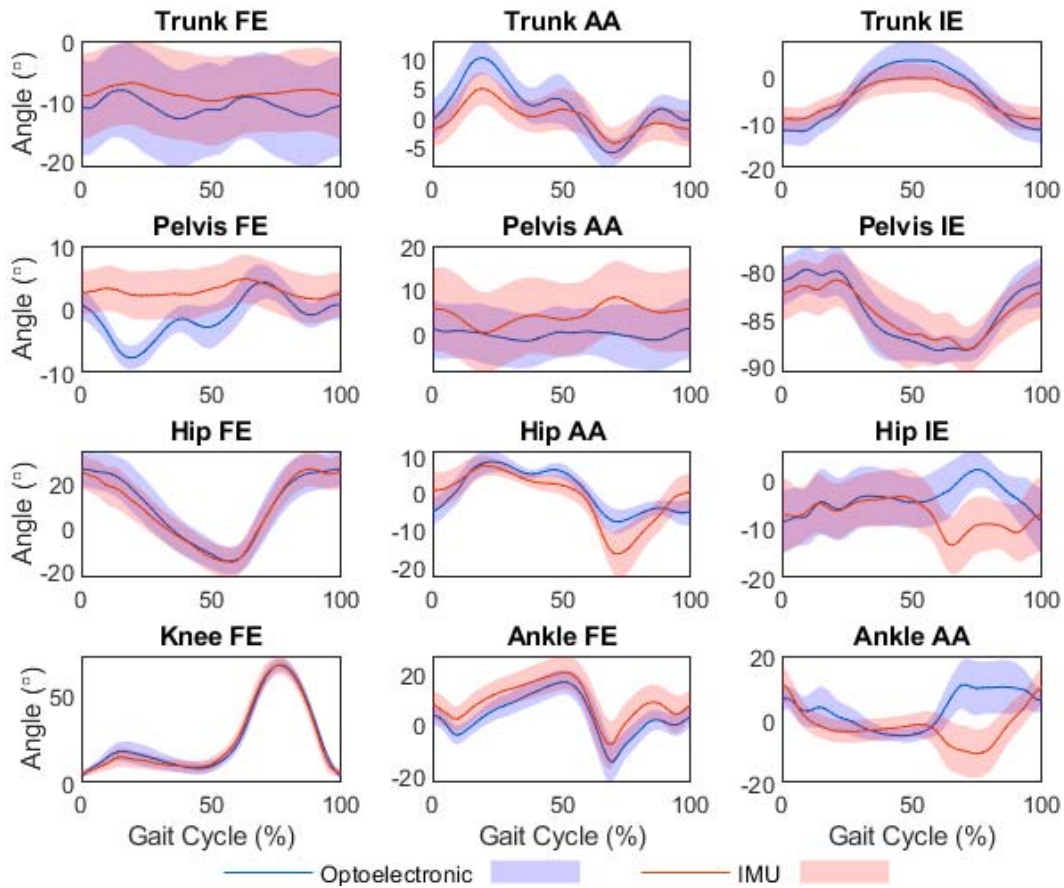
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Table 1. Root mean squared differences (RMSD) and RMSD relative to optoelectronic-modeled range of motion (CV_{rom}) of the IMU-modeled joint angle timeseries during gait ($N = 200$ strides). Values are group means [95% confidence intervals]. Highlights represent pooled differences accepted as reasonable (green: $RMSD \leq 5.0^\circ$), differences approaching reasonable levels (yellow: $5.0^\circ < RMSD \leq 7.0^\circ$), and differences exceeding reasonable levels (orange: $RMSD > 7.0^\circ$).

RMSD (°)	Angle	Gait condition					Pooled	
		Preferred speed, preferred swing	70% preferred speed, preferred swing	130% preferred speed, preferred swing	Preferred speed, active swing	Preferred speed, arms bound		
	Trunk FE	3.8 [2.1,5.5]	4.1 [2.1,5.5]	4.2 [2.2,6.2]	4.2 [2.8,5.6]	3.4 [2.0,4.8]	4.0 [2.3,5.6]	
	Trunk AA	3.6 [2.8,4.3]	3.6 [2.7,4.5]	3.9 [3.1,4.7]	4.3 [3.3,5.2]	3.4 [2.8,4.0]	3.7 [2.9,4.5]	
	Trunk IE	4.0 [3.1,5.0]	2.9 [2.5,3.3]	3.8 [3.2,4.4]	6.7 [1.9,11.4]	2.9 [2.5,3.4]	4.1 [2.6,5.5]	
	Pelvis FE	6.6 [5.4,7.7]	5.7 [4.4,7.0]	6.0 [5.0,7.0]	6.3 [4.9,7.6]	5.8 [4.4,7.3]	6.1 [4.8,7.3]	
	Pelvis AA	6.0 [4.7,7.3]	5.6 [4.4,6.8]	5.2 [3.9,6.4]	5.3 [4.2,6.5]	4.7 [3.4,6.1]	5.4 [4.1,6.6]	
	Pelvis IE	1.9 [1.5,2.2]	1.7 [1.3,2.2]	2.1 [1.7,2.5]	2.0 [1.6,2.4]	1.8 [1.5,2.2]	1.9 [1.5,2.3]	
	Hip FE	4.4 [3.5,5.2]	3.7 [3.0,4.4]	5.8 [4.9,6.8]	4.2 [3.4,4.9]	4.5 [3.0,6.0]	4.5 [3.6,5.5]	
	Hip AA	5.5 [4.6,6.4]	4.8 [4.0,5.6]	5.8 [4.9,6.7]	5.4 [4.4,6.4]	5.5 [4.6,6.5]	5.4 [4.5,6.3]	
	Hip IE	6.1 [5.4,6.8]	5.8 [4.9,6.6]	6.5 [5.7,7.2]	5.8 [5.0,6.7]	6.7 [5.5,7.9]	6.2 [5.3,7.0]	
	Knee FE	4.6 [3.7,5.5]	3.7 [3.1,4.3]	4.8 [3.6,5.9]	4.0 [3.6,4.4]	4.7 [3.7,5.6]	4.3 [3.5,5.1]	
	Ankle FE	6.7 [5.6,7.9]	6.2 [4.9,7.5]	7.5 [6.2,8.8]	6.9 [5.4,8.4]	7.4 [6.0,8.8]	6.9 [5.6,8.3]	
	Ankle AA	12.0 [10.3,13.6]	10.2 [8.8,11.5]	12.0 [10.7,13.2]	11.5 [10.1,12.9]	12.7 [10.9,14.5]	11.7 [10.2,13.2]	
	Pooled (all)	5.4 [3.7,7.1]	4.8 [3.3,6.3]	5.6 [4.0,7.3]	5.5 [3.5,7.6]	5.3 [3.4,7.2]	5.3 [3.6,7.1]	
	Pooled (without Ankle AA)	4.8 [4.4,5.2]	4.3 [4.0,4.6]	5.1 [4.6,5.5]	5.0 [4.5,5.5]	4.6 [4.2,5.0]	4.8 [4.4,5.2]	
CV _{rom} (%)	Trunk FE	25.5 [16.6,34.3]	27.9 [17.8,38.1]	29.4 [16.9,42.0]	27.7 [19.6,35.8]	28.2 [17.5,38.9]	27.7 [17.7,37.8]	
	Trunk AA	15.0 [13.0,17.1]	16.8 [13.7,20.0]	15.3 [13.0,17.7]	17.7 [14.5,20.8]	17.9 [15.1,19.2]	16.6 [13.9,19.2]	
	Trunk IE	15.6 [14.2,37.8]	14.2 [12.0,16.5]	13.7 [11.3,16.1]	22.1 [3.8,40.4]	15.6 [13.7,17.5]	16.2 [10.5,21.9]	
	Pelvis FE	40.7 [31.5,49.9]	39.1 [28.6,49.7]	33.4 [25.9,40.9]	35.6 [27.3,43.8]	43.3 [31.7,54.9]	38.4 [29.0,47.8]	
	Pelvis AA	67.9 [48.8,87.1]	56.4 [42.9,70.0]	59.1 [41.6,76.6]	53.7 [40.1,67.4]	57.6 [36.9,78.3]	59.0 [42.1,75.8]	
	Pelvis IE	13.4 [10.4,16.3]	12.9 [9.2,16.5]	12.8 [10.7,14.9]	11.7 [9.9,13.6]	13.2 [10.8,15.7]	12.8 [10.2,15.4]	
	Hip FE	9.1 [7.5,10.6]	8.5 [6.6,10.5]	11.0 [9.3,12.6]	8.0 [6.6,9.4]	9.2 [6.7,11.8]	9.2 [7.3,11.0]	
	Hip AA	23.3 [18.5,28.2]	21.9 [17.2,26.5]	22.9 [18.8,27.0]	22.1 [17.3,27.0]	28.2 [21.5,35.0]	23.7 [18.7,28.7]	
	Hip IE	32.2 [27.5,37.0]	30.2 [25.0,36.1]	32.9 [29.6,36.1]	29.8 [25.4,34.1]	35.0 [29.2,40.9]	32.0 [27.3,36.7]	
	Knee FE	6.7 [5.4,8.0]	5.5 [4.6,6.4]	7.0 [5.5,8.5]	6.0 [5.4,6.7]	6.8 [5.3,8.3]	6.4 [5.2,7.6]	
	Ankle FE	16.2 [13.9,18.5]	17.2 [14.3,20.2]	17.7 [15.2,20.2]	16.6 [13.3,19.9]	17.6 [14.5,20.6]	17.1 [14.3,19.9]	
	Ankle AA	46.1 [39.6,52.5]	41.4 [35.4,47.4]	46.1 [40.8,51.3]	43.5 [37.3,49.6]	49.8 [43.0,56.5]	45.3 [39.2,51.5]	
		Pooled (all)	26.0 [14.2,37.8]	24.3 [14.2,34.5]	25.1 [14.4,35.8]	24.6 [13.8,25.3]	26.9 [15.1,38.7]	25.4 [14.3,36.4]
		Pooled (without Ankle AA)	24.1 [21.0,27.3]	22.8 [20.2,25.4]	23.2 [20.5,25.9]	22.8 [19.7,26.0]	24.8 [21.5,28.1]	23.6 [20.6,26.5]

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FE: flexion/extension; AA: abduction/adduction; IE: internal/external rotation



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289 *Figure 2. Ensemble averaged joint angles from the optoelectronic and inertial measurement unit (IMU) models during gait at*
 290 *preferred speed and with preferred arm swing. Angles include flexion/extension (FE), abduction/adduction (AA), and*
 291 *internal/external rotation (IE).*

292 **3.2.2 Outcomes.** Mean values are presented in Table 2, with Bland-Altman plots for each

293 outcome displayed in Figures 3–7. Good–excellent consistency was seen for ROM of trunk IE,

294 pelvis IE, hip FE, and ankle FE ($ICC_{2,1}$: 0.76–0.85), for meanSD of trunk angles, pelvis IE, hip

295 angles, and ankle FE ($ICC_{2,1}$: 0.63–0.80), for λ_{max} of all angles except trunk IE and ankle AA

296 ($ICC_{2,1}$: 0.67–0.87), for DFA α of trunk IE and ankle FE ($ICC_{2,1}$: 0.62–0.65), and for SaEn of

297 trunk AA, pelvis IE, and knee FE ($ICC_{2,1}$: 0.61–0.74).

298 Differences [absolute: RMSD (relative: CV_{mean})] between outcomes with good–excellent

299 consistency were 2.0–7.7° (5.0–38.3%) for ROM, 0.14–0.61° (10.9–50.0%) for meanSD, 0.20–

300 0.72 (3.9–20.4%) for λ_{\max} , 0.16–0.17 (16.6–17.2%) for DFA α , and 0.07–0.24 (16.5–63.1%) for
301 SaEn.

302 Significant mean biases (defined as cases where the 95% confidence interval did not
303 cross zero; Table 2) indicated that the IMU model underestimated ROM of the trunk and sagittal
304 plane pelvis and ankle angles by 1.2–7.9°, but overestimated ROM of frontal plane pelvis, hip,
305 and ankle angles by 3.1–6.8° (Figure 3). For stride-to-stride variability outcomes, the IMU model
306 underestimated meanSD of trunk IE and pelvis AA by 0.26–0.52° (Figure 4), λ_{\max} of trunk IE by
307 0.23, DFA α of pelvis FE and hip AA by 0.09–0.11 (Figure 5), and SaEn of pelvis AA, hip AA,
308 hip IE, knee FE, and ankle AA by 0.02–0.34 (Figure 7). These outcomes were more frequently
309 overestimated by the IMU model, with higher meanSD (0.06–0.56°, Figure 4) and λ_{\max} (0.20–
310 0.65, Figure 5) for most angles, higher DFA α of ankle AA (0.12, Figure 6), and higher SaEn of
311 trunk AA, trunk IE, pelvis FE, pelvis IE, hip FE, and ankle FE (0.04–0.28, Figure 7). These
312 biases approximated the optoelectronic-measured inter-individual standard deviations in our
313 sample and in measurements from other studies for joint angle ROM [47], meanSD [33,37],
314 DFA α [37], and SaEn [5,37], but exceeded inter-individual standard deviations for joint angle
315 λ_{\max} [4,37]. The Bland-Altman plots (Figures 3–7) show that nearly all measurement differences
316 (14 participants \times 5 conditions = 70 values) fell within the 95% limits of agreement, with few
317 outliers in ROM (N = 1–5), meanSD (N = 2–5), λ_{\max} (N = 1–4), DFA α (N = 1–5), and SaEn (N =
318 0–6) of individual angles.

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Table 2. Validity of IMU-modeled vs. optoelectronic-modeled joint angle outcomes for gait. Outcomes are range of motion (ROM), mean standard deviation (meanSD), local divergence exponent (λ_{max}), detrended fluctuation analysis scaling exponent for range of motion (DFA α), and sample entropy (SaEn). Validity metrics are intraclass correlation coefficients ($ICC_{2,1}$), root mean square difference (RMSD), coefficient of variation of the optoelectronic mean (CV_{mean}), IMU-optoelectronic bias, and 95% limits of agreement ($LOA_{95\%}$). Values are group means [95% confidence intervals]. Highlights represent excellent (dark green: $ICC_{2,1} \geq 0.75$), good (green: $0.60 \leq ICC_{2,1} < 0.75$), fair (yellow: $0.40 \leq ICC_{2,1} < 0.60$), and poor (orange: $ICC_{2,1} < 0.40$) consistency. Asterisks identify significant bias (95% confidence interval did not cross zero).

	Angle	ICC _{2,1}	RMSD	CV _{mean}	Bias	LOA _{95%} Lower	Upper
ROM	Trunk FE	0.13 [-0.11,0.35]	2.5 [2.1,2.9]	26.1 [22.3,29.8]	-1.3 [-1.8,-0.8] *	-5.4 [-5.9,-4.9]	2.8 [2.3,3.3]
	Trunk AA	0.48 [0.28,0.64]	7.8 [7.0,8.6]	41.3 [38.3,44.4]	-7.0 [-7.8,-6.2] *	-13.6 [-14.3,-12.8]	-0.5 [-1.3,0.3]
	Trunk IE	0.85 [0.77,0.90]	7.7 [6.8,8.6]	38.3 [34.8,41.7]	-6.7 [-7.6,-5.8] *	-14.1 [-15.0,-13.2]	0.7 [-0.2,1.6]
	Pelvis FE	0.04 [-0.20,0.27]	8.5 [7.8,9.3]	60.5 [56.8,64.1]	-7.9 [-8.7,-7.1] *	-14.4 [-15.2,-13.6]	-1.4 [-2.2,-0.6]
	Pelvis AA	0.09 [-0.14,0.32]	4.1 [3.6,4.6]	85.1 [69.5,100.7]	3.3 [2.7,3.9] *	-1.5 [-2.0,-0.9]	8.1 [7.5,8.7]
	Pelvis IE	0.84 [0.75,0.90]	2.0 [1.7,2.4]	14.8 [12.1,17.6]	-0.5 [-0.9,0.0] *	-4.4 [-4.9,-3.9]	3.5 [3.0,4.0]
	Hip FE	0.84 [0.75,0.90]	2.8 [2.4,3.3]	5.0 [3.8,6.1]	0.1 [-0.6,0.8]	-5.5 [-6.2,-4.8]	5.7 [5.0,6.4]
	Hip AA	0.24 [-0.01,0.45]	8.5 [7.5,9.6]	44.8 [36.9,52.7]	6.8 [5.5,8.0] *	-3.5 [-4.9,-2.3]	17.1 [15.8,18.3]
	Hip IE	-0.11 [-0.35,-0.13]	4.8 [4.2,5.4]	36.9 [30.4,43.4]	0.4 [-0.7,1.6]	-9.1 [-10.2,-7.9]	9.9 [8.7,11.0]
	Knee FE	0.26 [0.02,0.47]	6.9 [5.9,7.9]	8.8 [7.2,10.3]	-1.4 [-3.0,0.2]	-14.7 [-16.4,13.1]	11.9 [10.3,13.5]
	Ankle FE	0.76 [0.64,0.85]	4.7 [4.1,5.4]	12.4 [10.3,14.5]	-1.2 [-2.3,-0.1] *	-10.3 [-11.4,-9.2]	7.8 [6.7,8.9]
	Ankle AA	0.10 [-0.14,0.34]	8.4 [7.2,9.6]	33.0 [26.6,39.3]	3.1 [1.2,5.0] *	-12.3 [-14.2,-10.4]	18.5 [16.6,20.3]
meanSD	Trunk FE	0.63 [0.46,0.75]	0.29 [0.25,0.34]	17.8 [14.1,21.6]	0.07 [0.00,0.14] *	-0.49 [-0.56,-0.42]	0.63 [0.56,0.70]
	Trunk AA	0.79 [0.68,0.86]	0.18 [0.15,0.22]	10.9 [8.6,13.1]	0.02 [-0.02,0.07]	-0.34 [-0.38,-0.29]	0.38 [0.34,0.43]
	Trunk IE	0.80 [0.70,0.87]	0.60 [0.54,0.66]	36.3 [33.6,39.1]	-0.52 [-0.59,-0.45] *	-1.10 [-1.17,-1.03]	0.06 [-0.01,0.13]
	Pelvis FE	0.54 [0.35,0.68]	0.29 [0.25,0.34]	34.7 [28.4,41.0]	0.22 [0.18,0.27] *	-0.16 [-0.21,-0.11]	0.60 [0.56,0.65]
	Pelvis AA	0.53 [0.34,0.68]	0.31 [0.27,0.34]	27.3 [24.4,30.2]	-0.26 [-0.30,-0.22] *	-0.58 [-0.61,-0.54]	0.05 [0.01,0.09]
	Pelvis IE	0.64 [0.48,0.76]	0.28 [0.23,0.32]	19.7 [15.1,24.2]	0.20 [0.15,0.24] *	-0.19 [-0.242,-0.15]	0.59 [0.54,0.63]
	Hip FE	0.80 [0.69,0.87]	0.23 [0.20,0.26]	17.5 [14.4,20.6]	0.16 [0.12,0.20] *	-0.16 [-0.20,-0.12]	0.48 [0.44,0.52]
	Hip AA	0.70 [0.56,0.81]	0.14 [0.12,0.16]	12.6 [10.1,15.0]	0.06 [0.03,0.09] *	-0.18 [-0.21,-0.15]	0.30 [0.27,0.33]
	Hip IE	0.69 [0.54,0.80]	0.20 [0.17,0.23]	12.5 [9.9,15.1]	0.10 [0.06,0.14] *	-0.24 [-0.28,-0.20]	0.44 [0.40,0.48]
	Knee FE	0.48 [0.27,0.65]	0.40 [0.34,0.46]	20.7 [15.6,25.9]	0.19 [0.10,0.27] *	-0.51 [-0.60,-0.43]	0.89 [0.80,0.97]
	Ankle FE	0.68 [0.53,0.79]	0.61 [0.55,0.66]	50.0 [43.5,56.5]	0.56 [0.50,0.62] *	0.10 [-0.04,0.16]	1.02 [0.97,1.08]
	Ankle AA	0.24 [0.00,0.46]	0.38 [0.32,0.43]	25.6 [20.7,30.5]	0.20 [0.13,0.28] *	-0.42 [-0.49,-0.34]	0.83 [0.75,0.90]
λ_{max}	Trunk FE	0.72 [0.58,0.82]	0.54 [0.48,0.60]	14.3 [12.3,16.2]	0.44 [0.37,0.52] *	-0.18 [-0.25,-0.10]	1.06 [0.99,1.14]
	Trunk AA	0.85 [0.77,0.91]	0.40 [0.36,0.44]	10.5 [9.3,11.7]	0.36 [0.32,0.40] *	0.04 [-0.00,0.08]	0.69 [0.65,0.72]
	Trunk IE	0.59 [0.40,0.73]	0.34 [0.29,0.39]	6.5 [5.3,7.7]	-0.23 [-0.29,-0.17] *	-0.73 [-0.79,-0.67]	0.27 [0.21,0.33]
	Pelvis FE	0.72 [0.57,0.82]	0.72 [0.65,0.78]	20.4 [18.2,22.5]	0.65 [0.58,0.73] *	0.07 [-0.00,0.14]	1.24 [1.17,1.31]
	Pelvis AA	0.79 [0.67,0.86]	0.34 [0.30,0.39]	9.1 [7.5,10.7]	0.24 [0.18,0.30] *	-0.25 [-0.31,-0.19]	0.73 [0.67,0.79]
	Pelvis IE	0.67 [0.51,0.78]	0.22 [0.19,0.25]	3.9 [3.1,4.7]	-0.04 [-0.09,0.01]	-0.47 [-0.52,-0.42]	0.39 [0.34,0.44]
	Hip FE	0.87 [0.80,0.92]	0.20 [0.17,0.22]	4.2 [3.5,4.9]	-0.04 [-0.09,0.01]	-0.42 [-0.47,-0.37]	0.34 [0.29,0.39]
	Hip AA	0.78 [0.66,0.86]	0.31 [0.28,0.34]	8.4 [7.5,9.4]	0.27 [0.23,0.30] *	-0.02 [-0.05,0.02]	0.56 [0.52,0.59]
	Hip IE	0.75 [0.62,0.84]	0.44 [0.39,0.49]	12.7 [10.9,14.5]	0.38 [0.33,0.43] *	-0.06 [-0.11,0.00]	0.82 [0.77,0.87]
	Knee FE	0.74 [0.61,0.84]	0.40 [0.34,0.46]	7.4 [5.9,8.9]	0.20 [0.12,0.28] *	-0.48 [-0.56,-0.40]	0.88 [0.79,0.96]
	Ankle FE	0.74 [0.60,0.83]	0.45 [0.41,0.50]	13.0 [11.4,14.5]	0.40 [0.35,0.45] *	-0.02 [-0.07,0.03]	0.82 [0.77,0.87]

DFAα	Ankle AA	0.37 [0.14,0.56]	1.17 [1.10,1.25]	51.1 [45.9,56.4]	1.13 [1.05,1.21] *	0.49 [0.41,0.56]	1.77 [1.69,1.85]
	Trunk FE	0.40 [0.18,0.58]	0.21 [0.18,0.24]	22.7 [17.2,28.2]	-0.03 [-0.08,0.02]	-0.44 [-0.49,-0.39]	0.38 [0.33,0.43]
	Trunk AA	0.27 [0.04,0.48]	0.19 [0.16,0.22]	18.7 [14.8,22.6]	-0.05 [-0.09,0.00]	-0.42 [-0.46,-0.37]	0.32 [0.28,0.37]
	Trunk IE	0.65 [0.49,0.76]	0.17 [0.15,0.20]	16.6 [13.0,20.2]	-0.00 [-0.04,0.04]	-0.34 [-0.38,-0.30]	0.34 [0.30,0.38]
	Pelvis FE	0.36 [0.14,0.54]	0.25 [0.21,0.29]	21.9 [17.9,25.8]	-0.11 [-0.16,-0.06] *	-0.55 [-0.60,-0.49]	0.33 [0.27,0.38]
	Pelvis AA	0.18 [-0.06,0.40]	0.23 [0.20,0.27]	28.6 [22.2,35.0]	0.04 [-0.01,0.10]	-0.41 [-0.47,-0.36]	0.50 [0.44,0.55]
	Pelvis IE	0.46 [0.25,0.63]	0.16 [0.14,0.19]	17.6 [13.3,21.9]	0.03 [-0.01,0.06]	-0.29 [-0.33,-0.25]	0.34 [0.30,0.38]
	Hip FE	0.50 [0.30,0.66]	0.20 [0.17,0.23]	21.0 [16.8,25.3]	-0.02 [-0.06,0.03]	-0.41 [-0.46,-0.36]	0.38 [0.33,0.43]
	Hip AA	0.37 [0.14,0.56]	0.23 [0.20,0.27]	21.3 [17.4,25.1]	-0.07 [-0.12,-0.01] *	-0.50 [-0.56,-0.45]	0.37 [0.32,0.43]
	Hip IE	0.13 [-0.12,0.36]	0.24 [0.21,0.27]	26.7 [21.8,31.6]	-0.04 [-0.10,0.02]	-0.51 [-0.56,-0.45]	0.43 [0.37,0.48]
	Knee FE	0.43 [0.21,0.61]	0.20 [0.17,0.23]	20.0 [16.1,23.9]	0.02 [-0.03,0.07]	-0.37 [-0.42,-0.32]	0.41 [0.36,0.46]
	Ankle FE	0.62 [0.44,0.75]	0.16 [0.13,0.18]	17.2 [13.4,21.0]	0.02 [-0.02,0.06]	-0.29 [-0.33,-0.25]	0.33 [0.29,0.37]
SaEn	Ankle AA	-0.05 [-0.29,0.20]	0.31 [0.35,0.26]	35.9 [28.2,43.7]	0.12 [0.05,0.19] *	-0.44 [-0.51,-0.38]	0.68 [0.61,0.75]
	Trunk FE	0.56 [0.38,0.70]	0.18 [0.15,0.20]	17.0 [13.8,20.2]	0.02 [-0.02,0.06]	-0.33 [-0.37,-0.29]	0.36 [0.32,0.41]
	Trunk AA	0.61 [0.44,0.74]	0.19 [0.16,0.21]	29.8 [23.3,36.3]	0.15 [0.13,0.18] *	-0.05 [-0.08,-0.03]	0.36 [0.34,0.39]
	Trunk IE	0.48 [0.28,0.64]	0.10 [0.08,0.12]	16.1 [12.3,19.9]	0.04 [0.01,0.06] *	-0.15 [-0.17,-0.13]	0.22 [0.20,0.24]
	Pelvis FE	0.05 [-0.19,0.28]	0.33 [0.29,0.37]	49.2 [40.8,57.6]	0.28 [0.23,0.32] *	-0.09 [-0.13,-0.04]	0.64 [0.59,0.68]
	Pelvis AA	0.22 [-0.01,0.43]	0.41 [0.36,0.47]	31.6 [28.1,35.2]	-0.34 [-0.39,-0.28] *	-0.82 [-0.87,-0.76]	0.14 [0.09,0.20]
	Pelvis IE	0.64 [0.48,0.76]	0.24 [0.21,0.27]	63.1 [47.9,78.4]	0.19 [0.15,0.22] *	-0.11 [-0.14,-0.07]	0.48 [0.45,0.52]
	Hip FE	0.44 [0.23,0.62]	0.05 [0.04,0.06]	17.9 [14.3,21.4]	0.04 [0.03,0.04] *	-0.04 [-0.05,-0.03]	0.11 [0.10,0.12]
	Hip AA	0.17 [-0.08,0.39]	0.15 [0.13,0.17]	23.5 [20.0,27.0]	-0.09 [-0.12,-0.06] *	-0.33 [-0.36,-0.30]	0.15 [0.12,0.18]
	Hip IE	-0.12 [-0.35,0.12]	0.25 [0.22,0.29]	21.3 [17.2,25.4]	-0.04 [-0.10,-0.02] *	-0.53 [-0.59,-0.47]	0.46 [0.40,0.52]
	Knee FE	0.74 [0.61,0.83]	0.07 [0.05,0.08]	16.5 [12.8,20.2]	-0.02 [-0.04,-0.01] *	-0.14 [-0.16,-0.13]	0.10 [0.08,0.11]
	Ankle FE	0.46 [0.24,0.63]	0.13 [0.11,0.14]	24.6 [20.2,29.0]	0.10 [0.08,0.12] *	-0.06 [-0.08,-0.04]	0.26 [0.24,0.28]
Ankle AA	0.10 [-0.14,0.34]	0.19 [0.16,0.22]	26.8 [22.1,31.6]	-0.03 [-0.08,-0.01] *	-0.40 [-0.44,-0.35]	0.34 [0.29,0.38]	

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FE: flexion/extension; AA: abduction/adduction; IE: internal/external rotation

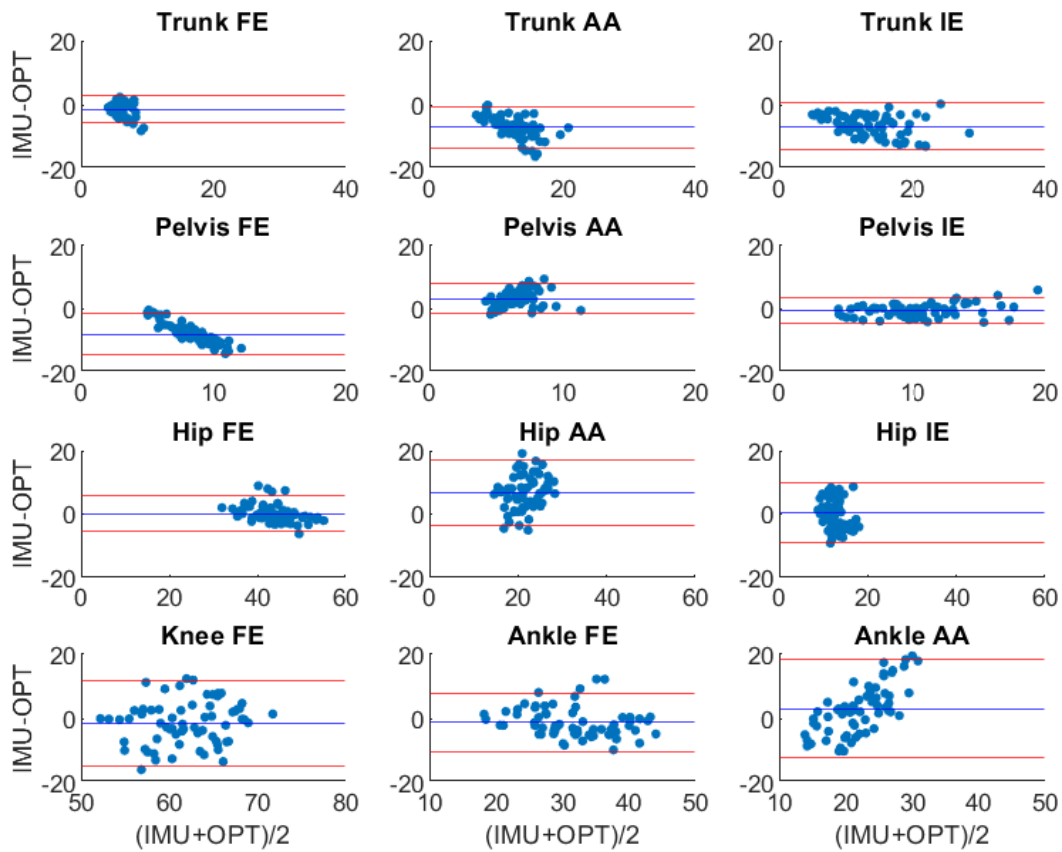


Figure 3. Bland-Altman plots of *range of motion* (ROM) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (OPT) and IMU-driven biomechanical models of constant-speed treadmill gait. Mean bias (blue line) and 95% limits of agreement (red lines) are shown. Values are in degrees ($^{\circ}$). Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

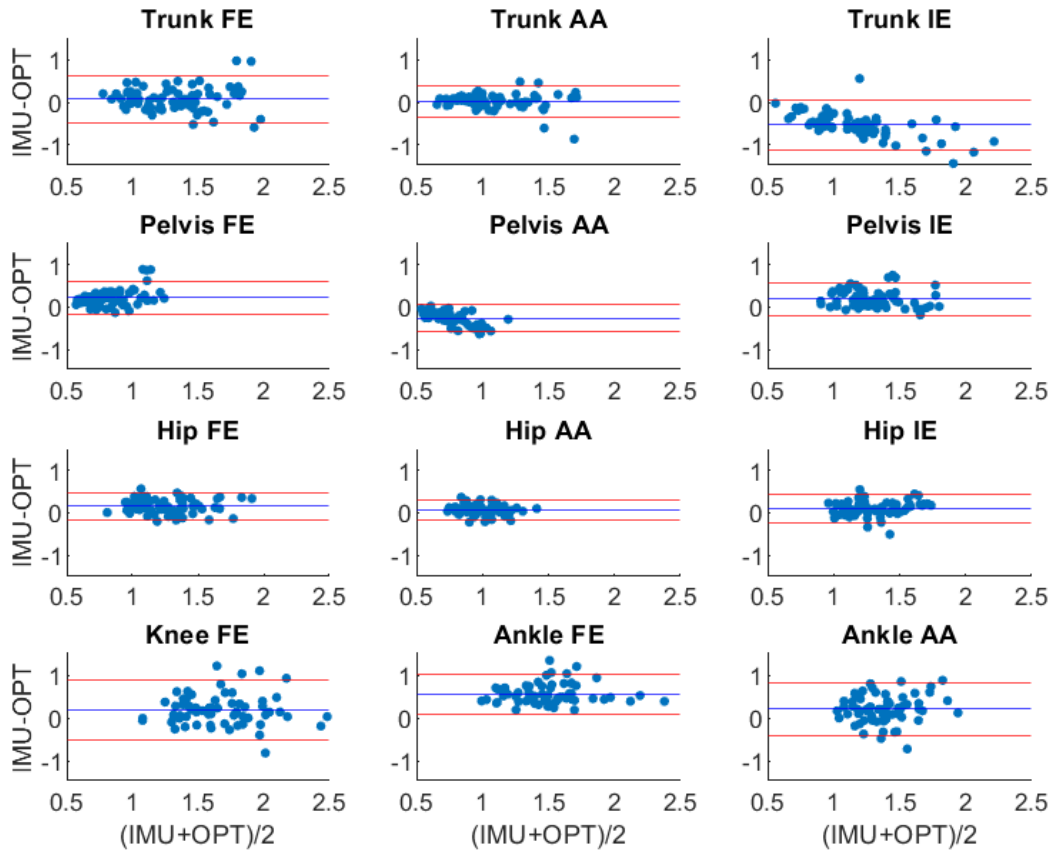


Figure 4. Bland-Altman plots of mean standard deviation (meanSD) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (OPT) and IMU-driven biomechanical models of constant-speed treadmill gait. Mean bias (blue line) and 95% limits of agreement (red lines) are shown. Values are in degrees ($^{\circ}$). Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

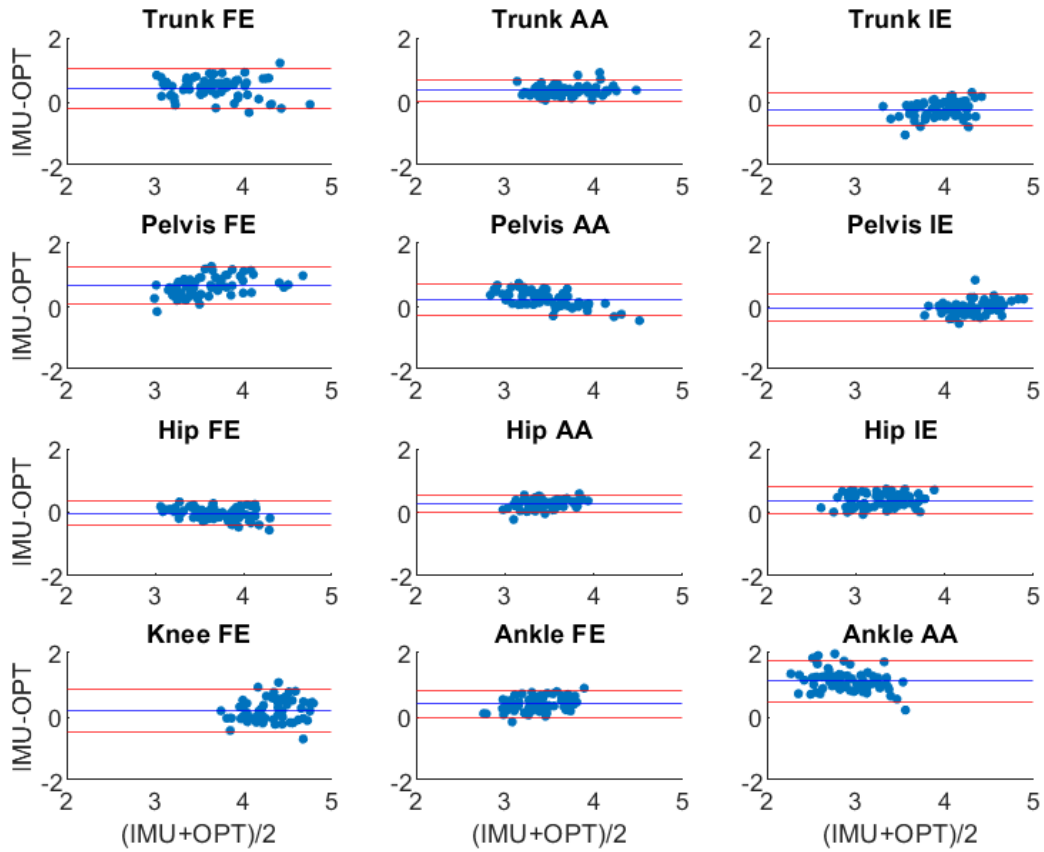


Figure 5. Bland-Altman plots of local divergence exponent (λ_{max}) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (OPT) and IMU-driven biomechanical models of constant-speed treadmill gait. Mean bias (blue line) and 95% limits of agreement (red lines) are shown. Values are in arbitrary units. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

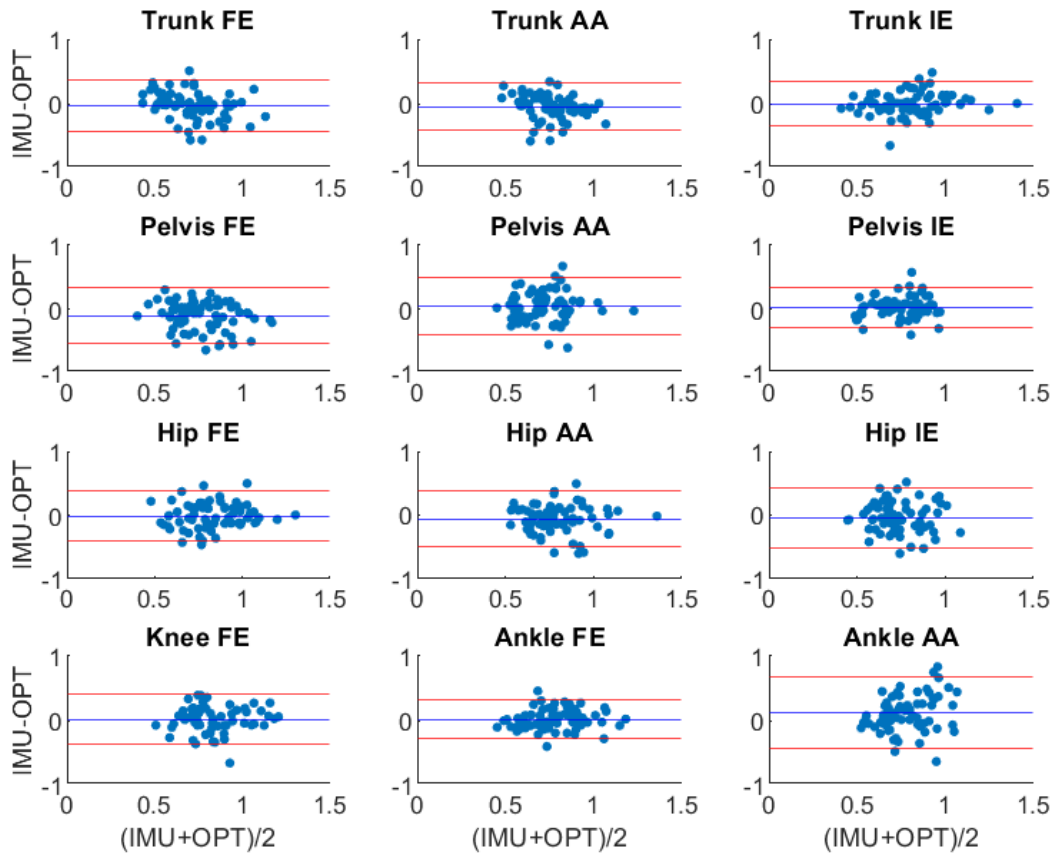


Figure 6. Bland-Altman plots of *detrended fluctuation analysis scaling exponent (DFA α)* of range of motion of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (OPT) and IMU-driven biomechanical models of constant-speed treadmill gait. Mean bias (blue line) and 95% limits of agreement (red lines) are shown. Values are in arbitrary units. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

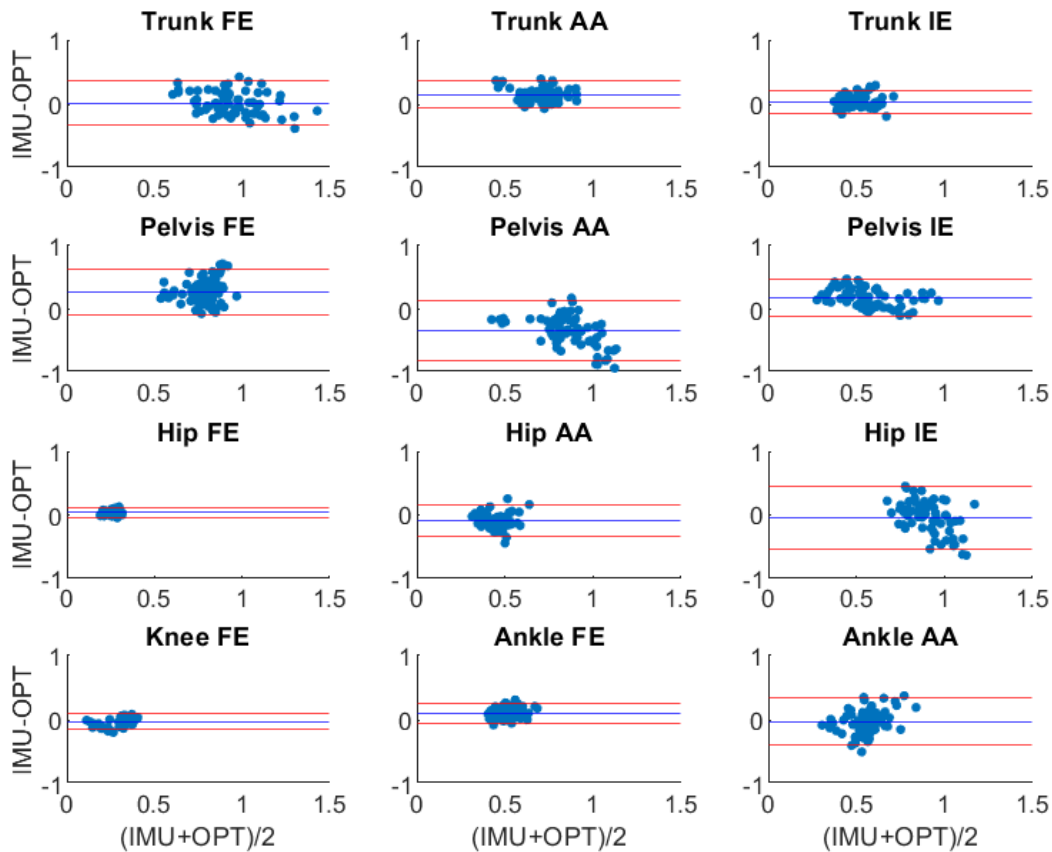


Figure 7. Bland-Altman plots of *sample entropy* (*SaEn*) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (OPT) and IMU-driven biomechanical models of constant-speed treadmill gait. Mean bias (blue line) and 95% limits of agreement (red lines) are shown. Values are in arbitrary units. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

3.2 IMU-model sensitivity

Precise descriptives (means, 95% confidence intervals) and statistical effects (p-values) for each outcome can be found in Supplementary material (Tables S1 and S2). Since consistency was generally poor–moderate for DFA α and SaEn, indicating a lack of acceptable concurrent validity, the sensitivity of these outcomes was not explored.

ROM (Figure 8). *Speed effects:* The models detected similar responses relative to preferred-speed gait, with decreased ROM of trunk AA, trunk IE, hip FE, hip IE, and ankle FE at

70% preferred speed, and increased ROM of trunk AA, trunk IE, pelvis IE, hip angles, and ankle FE at 130% preferred speed. Differing responses, where only one model detected a significant change, were found at 70% preferred speed for pelvis angles, hip AA, knee FE, and ankle AA, and at 130% preferred speed for pelvis FE, pelvis AA, and ankle AA. The changes detected, however, followed the same trends of decreased ROM at 70% preferred speed and increased ROM at 130% preferred speed. *Swing effects:* The models also detected similar changes relative to preferred arm swing during active swing, with increased ROM of pelvis IE and hip FE, decreased ROM of knee FE, and no change in ROM of trunk FE, trunk AA, pelvis FE, pelvis AA, hip AA, hip IE, ankle FE, and ankle AA. With arms bound, both models detected decreased ROM of trunk IE and no change for hip FE, hip IE, ankle FE, and ankle AA.

meanSD (Figure 9). *Speed effects:* The models detected similar non-responses to speed changes for meanSD of trunk FE, trunk IE, ankle FE, and ankle AA. However, optoelectronic-based model increases in meanSD at 70% preferred speed (trunk AA, pelvis FE, pelvis AA, hip FE, hip AA, knee FE) and at 130% preferred speed (pelvis IE) went undetected by the IMU-based model. *Swing effects:* Relative to preferred arm swing, the models detected similar increases in meanSD (trunk AA, pelvis AA, pelvis IE, hip AA, hip IE) and non-responses (trunk FE, hip FE, knee FE) during active arm swing, as well as similar non-responses with arms bound (trunk FE, pelvis FE, pelvis AA, and all hip, knee, and ankle angles). Optoelectronic-detected increases in meanSD of trunk AA and IE and decreases for pelvis IE with arms bound went undetected by the IMU-based model.

λ_{\max} (Figure 10). *Speed effects:* The models detected similar responses to speed changes in λ_{\max} of joint angles with few exceptions. Both models detected increases in λ_{\max} at 70% preferred speed (trunk FE, trunk AA, pelvis-down angles), decreases in λ_{\max} at 130% preferred

speed (trunk FE, pelvis FE, pelvis AA, hip AA, hip IE, ankle AA), and no response in λ_{\max} of knee FE at 130% preferred speed. *Swing effects*: The models also detected similar λ_{\max} responses relative to preferred arm swing, with increases in trunk FE, pelvis angles, hip AA, hip IE, and ankle FE during active swing, and decreases in trunk IE, but increases in pelvis IE, with arms bound. Similar non-responses in λ_{\max} of ankle AA during active swing and of trunk FE, hip FE, hip AA, knee FE, ankle FE, and ankle AA with arms bound were also seen.

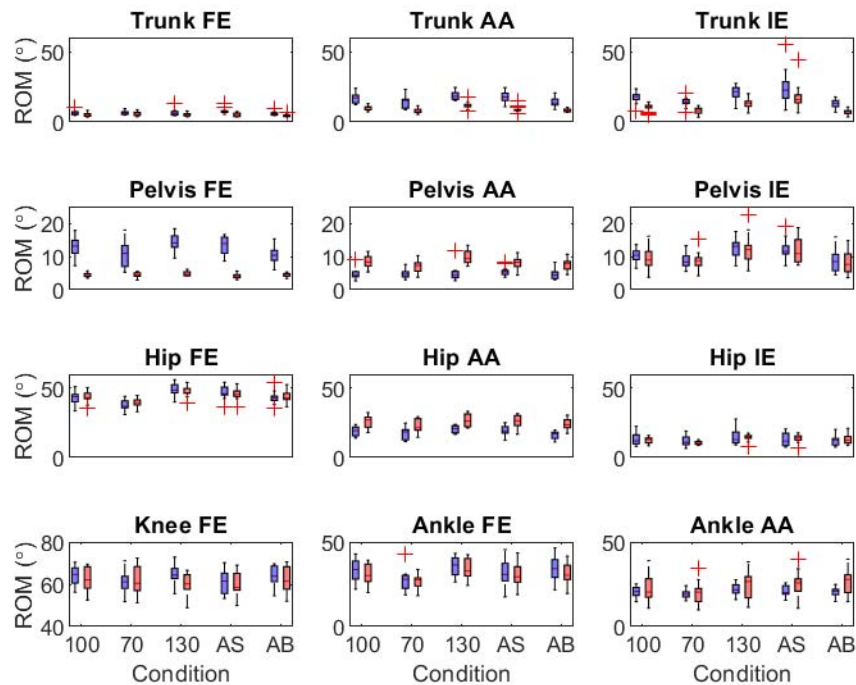


Figure 8. Boxplots of range of motion (ROM) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (blue) and IMU-driven (red) biomechanical models of constant-speed treadmill gait. Conditions displayed on the horizontal axis are preferred speed and arm swing (100), 70% preferred speed (70), 130% preferred speed (130), active arm swing (AS), and arms bound (AB). Red crosses are outliers $> 2 * \text{interquartile range}$. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

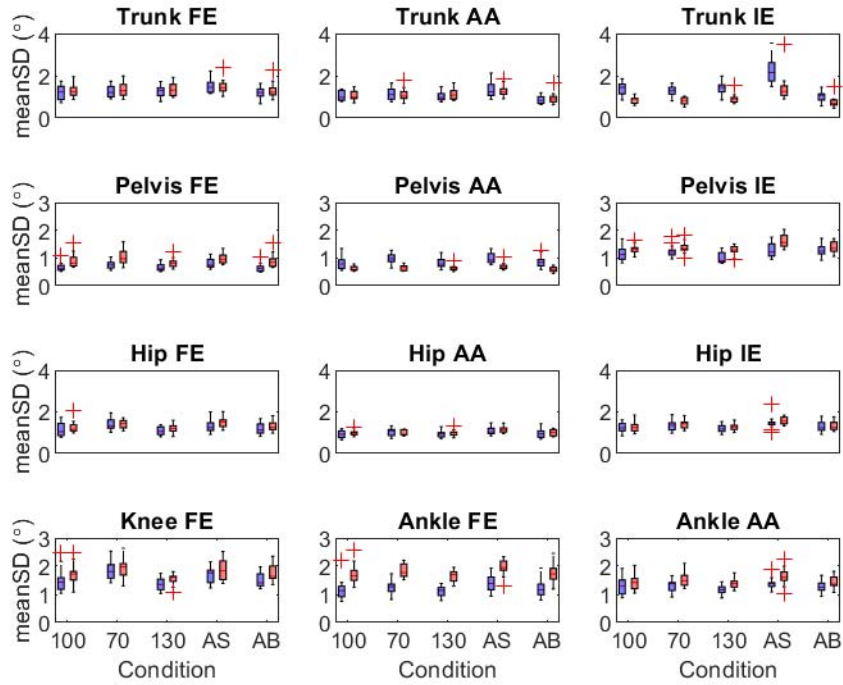


Figure 9. Boxplots of mean standard deviation (*meanSD*) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (blue) and IMU-driven (red) biomechanical models of constant-speed treadmill gait. Conditions displayed on the horizontal axis are preferred speed and arm swing (100), 70% preferred speed (70), 130% preferred speed (130), active arm swing (AS), and arms bound (AB). Red crosses are outliers $> 2 * \text{interquartile range}$. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

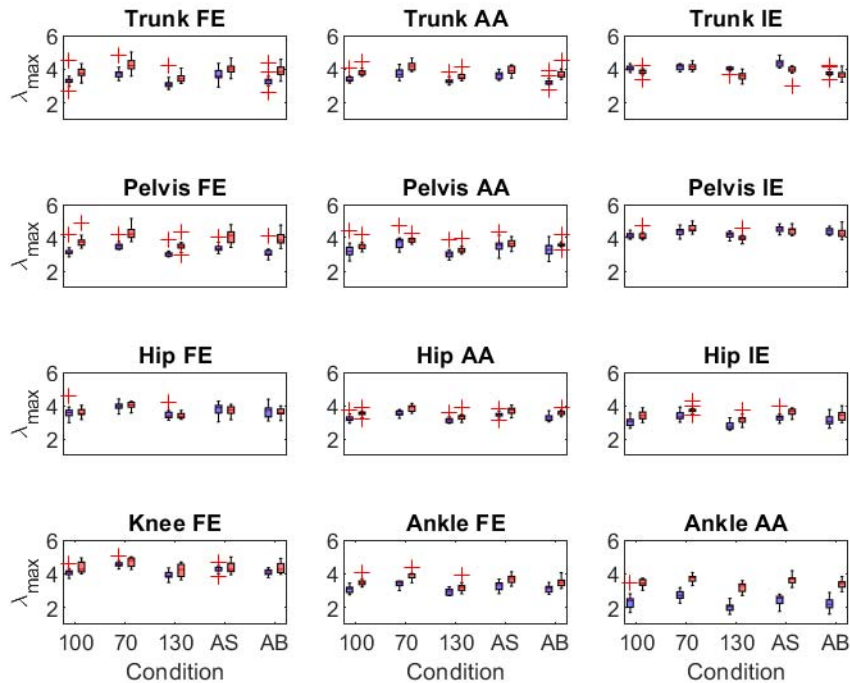


Figure 10. Boxplots of local divergence exponent (λ_{max}) of trunk, pelvis, and lower-limb joint angles for the optoelectronic-driven (blue) and IMU-driven (red) biomechanical models of constant-speed treadmill gait. Conditions displayed on the horizontal axis are preferred speed and arm swing (100), 70% preferred speed (70), 130% preferred speed (130), active arm swing (AS), and arms bound (AB). Red crosses are outliers $> 2 * \text{interquartile range}$. Angles include flexion/extension (FE), abduction/adduction (AA), and internal/external rotation (IE).

4. Discussion

4.1 Validity of IMU-modeled joint angle timeseries

Using a full-body biomechanical model with muscle-actuated lower limbs [31] to analyze several 7-minute conditions of gait kinematics, our findings confirm that driving this model with IMUs (using the *OpenSense* open-source toolkit for *OpenSim*) produces accurate joint angle timeseries relative to the optoelectronic-driven model, with the exception of ankle inversion. RMSD across all gait speed and arm swing conditions averaged 5.3° ; when ankle inversion was excluded from this calculation, RMSD averaged 4.8° , an accuracy viewed as acceptable for many clinical applications [27]. This amount of accuracy and the range in RMSD among

individual joints (1.7–7.5°) agrees with recent findings from the developers [13,14]. One observation from our ensemble-averaged lower-limb joint angles (Figure 2) was that optoelectronic–IMU differences were most notable in the non-sagittal plane during swing (e.g., hip abduction, hip rotation, and ankle inversion), potentially due to the relatively small ROMs of these angles in combination with the higher segmental velocities during swing. Nonetheless, we report an accuracy for hip rotation (pooled RMSD of 6.2°) that improved upon the developers’ findings (10–12°). Because allowing ankle inversion/eversion in addition to dorsiflexion/plantarflexion did not affect the inverse kinematics solution for hip rotation (see, for example, Figure S1), we believe our improvement in accuracy is attributable to omitting magnetometer data when using Madgwick’s gradient-descent fusion algorithm [19] and, instead, identifying negative (< -0.0010 rad/s) and positive (> 0.0010 rad/s) linear orientation drifts and detrending these offline. Extending previous findings, we also found RMSDs of IMU-modeled trunk and pelvis angles (1.7–6.6°) that were similar to those of the joints in the lower limb. Thus, with the notable exception of ankle inversion/eversion, our findings indicate that joint angle timeseries for the trunk and below can be estimated on average with acceptable accuracy during walking over a wide range of walking speeds and arm swing amplitudes.

4.2 Validity of IMU-modeled joint angle outcomes

The concurrent validity of the joint angle timeseries, however, did not extend to the discrete outcomes to the same extent. For ROM, as reported by Beange et al. [48,49], optoelectronic–IMU consistency was excellent in the primary plane of movement (trunk rotation, pelvis rotation, hip flexion, ankle dorsiflexion) with RMSDs similar to the corresponding timeseries (ROM: 2.0–7.7°; timeseries: 1.9–6.9°), but consistency was often poor in the non-primary movement planes. Therefore, measurement consistency of ROM for the secondary and

tertiary planes was not improved by using a biomechanically constrained kinematic model. RMSDs for the secondary and tertiary planes (2.5–8.5°) were comparable to those of the primary movement plane, confirming the findings of Beange et al. [48,49] and likely due to the lower ROMs about the secondary and tertiary planes; future IMU-modelling work of lower-limb joint kinematics could investigate whether consistency is higher in locomotor tasks with similar segmental velocities that include larger ROMs in the frontal and transverse planes, such as walking with side-stepping and/or turning. ROM was also below proposed limits of agreement of $\pm 10^\circ$ [50] for only 5 of the 12 degrees of freedom we analyzed, with underestimation biases (trunk rotation and ankle flexion) partly responsible. Together, these findings show that acceptable validity of a joint angle timeseries does not necessarily translate to acceptable validity in the discrete outcomes of that timeseries. Specifically, our IMU model estimates joint angle ROM accurately, but has inconsistencies in the non-primary movement planes and limitations in absolute agreement.

For motor variability outcomes, consistency was best for λ_{\max} and meanSD, reaching good–excellent levels for nearly all degrees of freedom with RMSDs of 0.22–0.72° and 0.14–0.61°, respectively. Our findings agree with the high optoelectronic–IMU consistency in λ_{\max} reported for repetitive spine flexion/extension [51] and support the concurrent validity of IMU-based measurements of joint local dynamic stability. For the first time, we demonstrate high consistency and accuracy for joint angles with low range of motion and beyond the primary plane of movement, providing a new validated model for investigating magnitude of variability and local dynamic stability in tridimensional joint motion from stride to stride. Caution is advised regarding the absolute agreement of our outcomes since we were unable to identify limits of agreement in the literature for comparison and since the IMU-based model biases

showed that λ_{\max} and meanSD were each underestimated for trunk rotation but overestimated for hip, knee, and ankle angles.

DFA α and SaEn were not sufficiently consistent to be considered valid in their present computational form (DFA α ICC $_{2,1}$: -0.05–0.65; SaEn ICC $_{2,1}$: -0.12–0.74). DFA α inconsistencies may be partly derived from ROM inconsistencies since we investigated fluctuations of this discrete metric. This is supported by our finding that the only two angles with good consistency in DFA α (trunk rotation, ankle dorsiflexion) also had excellent consistency in ROM, indicating that improvement in calculation of ROM is likely to produce more valid analyses of fluctuation persistence. SaEn inconsistencies may be due to changes in the information of joint angles in the IMU-driven model. The calculation of this metric for continuous series of joint angles is sensitive to the sampling frequency, the tolerance ratio r , and the embedding dimension m [44]. We resampled IMU-based joint angles to compensate for sampling frequency and selected r and m values which produced stable measurements in slow and fast gait [43]. However, specific tuning of SaEn parameters to IMU-based kinematic models is likely needed. Alternatively, calculation of SaEn may be more robust to tuning parameters by analyzing the discrete ROM series rather than the continuous joint angle series [44], but this also relies on improving the estimation of ROM. Therefore, use of IMU-based biomechanical models to investigate the persistence and regularity of joint angle fluctuations in gait requires improvements to the estimation of ROM and the identification of appropriate SaEn calculation parameters for IMU-derived joint angles.

4.3 Sensitivity of IMU-modeled joint angle outcomes to within-participant effects

Our results show, for the first time, that joint angle outcomes estimated from an IMU-driven biomechanical model were also sensitive to within-participant responses detected by an optoelectronic-driven model in the majority of cases. Relative to gait at preferred speed and with preferred arm swing, responses to changes in speed and arm swing amplitude were detected identically between models for 28/48 comparisons for ROM, 29/48 comparisons for meanSD, and 35/48 comparisons for λ_{\max} . Model-detected responses are in agreement with several previous findings, including increases in joint ROM with fast speed and decreases with slow speed [52], increases in SD of trunk kinematics and meanSD of lower-limb angles with active arm swing [36,37], and increases in λ_{\max} of hip abduction with active arm swing [37]. Although our trunk motion responses appear to disagree with reported increases in meanSD with fast speed [33], increases in λ_{\max} with fast speed [34], and decreases in λ_{\max} with active arm swing [35], differences can be attributed to the reference frame of the trunk (relative to the pelvis in our study vs. relative to ground in [33]) and to the sensitivity of the λ_{\max} state-space to different inputs [53] (time-delayed joint angles in our study vs. velocities and accelerations [34] vs. time-delayed velocities [35]). In the minority of cases where our optoelectronic and IMU models detected different responses, directionality never disagreed, and differences were mostly due to the IMU-based model detecting no response when a change was detected by the optoelectronic-based model. In these cases, there were no statistically significant changes for the IMU-based model, but all means moved in the same direction as the optoelectronic-based model (Tables S1 and S2). As the probability of this due to chance is very low, it seems that IMU-modeled responses had smaller effect sizes than this study was powered to detect, due to smaller changes in magnitude and/or larger group variance. Furthermore, the excess limits of agreement for some ROMs did not influence sensitivity of the IMU model, as responses in ROM for angles

exceeding limits of $\pm 10^\circ$ [50] were the same in many cases during gait at fast speed (trunk abduction, trunk rotation, ankle dorsiflexion), at slow speed (trunk abduction, trunk rotation, hip abduction, ankle dorsiflexion), with active arm swing (trunk abduction, pelvis flexion, hip abduction, knee flexion, ankle dorsiflexion, ankle inversion), and with bound arms (trunk rotation, ankle flexion, ankle inversion). Together, these findings show that the IMU-driven biomechanical model is sensitive to within-group responses in ROM, meanSD, and λ_{\max} , and suggest that larger sample sizes may further improve sensitivity by compensating for smaller effect sizes.

4.4 Limitations

IMU-driven biomechanical simulations of movement are a new area for investigation and have important limitations. In our study, these were specific to the demographics of our sample, the biomechanical model, and how gait was studied. As a first step for modelling motor variability in gait, we recruited a convenience sample of healthy young adult males and females whose average BMI was in the “normal” range. Our motivation for investigating motor variability in gait, however, is to investigate stride-to-stride control of older adults as it relates to fall risk [3]. Confirmation of validity and sensitivity of an IMU-driven biomechanical model is still needed in older populations living with and without neurological conditions.

To mitigate differences attributed to different initial model poses in our evaluation of validity, we decided to offset the optoelectronic model to match the standing pose of the IMU model at the first frame, similar to the approach of Al Borno et al. [14]. This is unrealistic for situations where an optoelectronic system is unavailable to establish the initial IMU model pose. Adding the average correction (3.8° [14]) to the RMSD differences would produce differences

that exceed limits tolerable for clinical applications [27]. However, ROM and motor variability were unaffected by this offset, and further IMU calibration procedures [54] are unlikely to affect validity or sensitivity of these measures. Thus, we expect that our conclusions on the validity of our outcomes can be generalized to situations where only IMU sensors are available.

Finally, we investigated performance of the IMU-driven model during treadmill gait, where stride-to-stride variability is lower than in overground gait [55,56]. Treadmills allow for analyses at constant speed and over a large number of continuous strides; for example, more than 100 strides are needed to reliably quantify SD of stride time and λ_{\max} of tridimensional trunk acceleration [57]. Large numbers of strides can be recorded in optoelectronic-based overground studies when gait involves turning and strides are discontinuous [58] but calculation of local dynamic stability requires continuous timeseries. Suitability of IMU-driven biomechanical models for joint kinematic evaluations in overground locomotion is promising [13,14,25] but should be confirmed for evaluations of stride-to-stride control.

5. Conclusion

In summary, excluding ankle inversion, IMU-based joint angle timeseries were acceptably accurate from the trunk down, ROM was acceptably consistent and accurate in the primary plane of motion, and magnitude of variability and local dynamic stability were acceptably consistent and accurate in all planes of motion. Validity was supported by the sensitivity of the IMU model to gait speed and arm swing amplitude-related responses in ROM, magnitude of variability, and local dynamic stability in the majority of cases. However, IMU-modeled estimates of ROM fluctuation persistence and of angle regularity were not acceptably consistent or accurate. We conclude that, for moderate-duration walking at slow and fast speeds,

the IMU-driven, magnetometer-free, open-source biomechanical model used in this study provides valid estimates of joint kinematics, stride-to-stride magnitude of variability, and stride-to-stride local dynamic stability which are also sensitive to within-participant responses. This provides a new way to evaluate biomechanical control of walking outside of the lab and clinic, a step towards better predicting falls in aging adults.

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Author Contributions

Conceptualization, Christopher Bailey, Thomas Uchida, Julie Nantel and Ryan Graham; Data curation, Christopher Bailey; Formal analysis, Christopher Bailey, Thomas Uchida, Julie Nantel and Ryan Graham; Funding acquisition, Christopher Bailey, Julie Nantel and Ryan Graham; Investigation, Christopher Bailey; Methodology, Christopher Bailey, Thomas Uchida, Julie Nantel and Ryan Graham; Project administration, Christopher Bailey, Julie Nantel and Ryan Graham; Software, Christopher Bailey; Supervision, Julie Nantel and Ryan Graham; Validation, Christopher Bailey, Thomas Uchida, Julie Nantel and Ryan Graham; Visualization, Christopher Bailey; Writing – original draft, Christopher Bailey; Writing – review & editing, Christopher Bailey, Thomas Uchida, Julie Nantel and Ryan Graham.

Institutional Review Board Statement

The study was conducted according to the guidelines of the Declaration of Helsinki, and approved by the institutional Research Ethics Board of the University of Ottawa (H-01-21-6261).

Informed Consent Statement

Informed consent was obtained from all participants involved in the study.

Data Availability Statement

The data underlying the findings of this study will be made freely available upon publication.

Conflicts of Interest

The authors declare no conflicts of interest.

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