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# Complexity of human walking: the attractor complexity index is sensitive to gait synchronization with visual and auditory cues

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**Background.** During steady walking, gait parameters fluctuate from one stride to another with complex fractal patterns and long-range statistical persistence. When a metronome is used to pace the gait (sensorimotor synchronization), long-range persistence is replaced by stochastic oscillations (anti-persistence). Fractal patterns present in gait fluctuations are most often analyzed using detrended fluctuation analysis (DFA). This method requires the use of a discrete times series, such as intervals between consecutive heel strikes, as an input. Recently, a new nonlinear method, the attractor complexity index (ACI), has been shown to respond to complexity changes like DFA. But in contrast to DFA, ACI can be applied to continuous signals, such as body accelerations. The aim of this study was to further compare DFA and ACI in a treadmill experiment that induced complexity changes through sensorimotor synchronization. **Methods.** Thirty-six healthy adults walked 30 minutes on an instrumented treadmill under three conditions: no cueing, auditory cueing (metronome walking), and visual cueing (stepping stones). The center-of-pressure trajectory was discretized into time series of gait parameters, after which a complexity index (scaling exponent alpha) was computed via DFA. Continuous pressure position signals were used to compute the ACI. Correlations between ACI and DFA were then analyzed. The predictive ability of DFA and ACI to differentiate between cueing and no-cueing conditions was assessed using regularized logistic regressions and areas under the receiver operating characteristic curves (AUROC). **Results.** DFA and ACI were both significantly different among the cueing conditions. DFA and ACI were correlated (Pearson's  $r = 0.78$ ). Logistic regressions showed that DFA and ACI could differentiate between cueing/no cueing conditions with a high degree of confidence (AUROC = 1.0 and 0.96, respectively). **Conclusion.** Both DFA and ACI responded similarly to changes in cueing conditions and had comparable predictive power. This support the assumption that

ACI could be used instead of DFA to assess the long-range complexity of continuous gait signals.

1 **Complexity of human walking: the attractor**  
2 **complexity index is sensitive to gait synchronization**  
3 **with visual and auditory cues**

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19

## 20 **Abstract**

21 **Background.** During steady walking, gait parameters fluctuate from one stride to  
22 another with complex fractal patterns and long-range statistical persistence. When a  
23 metronome is used to pace the gait (sensorimotor synchronization), long-range  
24 persistence is replaced by stochastic oscillations (anti-persistence). Fractal patterns  
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29 DFA. But in contrast to DFA, ACI can be applied to continuous signals, such as body  
30 accelerations. The aim of this study was to further compare DFA and ACI in a treadmill  
31 experiment that induced complexity changes through sensorimotor synchronization.

32 **Methods.** Thirty-six healthy adults walked 30 minutes on an instrumented treadmill  
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37 ACI. Correlations between ACI and DFA were then analyzed. The predictive ability of  
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47 used instead of DFA to assess the long-range complexity of continuous gait signals.  
48

## 49 **Introduction**

50 Gait is a stereotyped sequence of movements that enable human beings to move through their  
51 environment. A fluid and stable gait requires the complex coordination of dozens of muscles  
52 controlling multiple joints. Biomechanical and energy constraints limit the range of gait  
53 movements to a narrow window (Holt et al., 1995); for example, at a preferred walking speed,  
54 step length and step time vary by only a few percent (Terrier, Turner & Schutz, 2005). It was  
55 previously thought that these small variations were random noise introduced by residual  
56 neuromuscular inaccuracies; however, after studying the structure of gait variability among  
57 hundreds of consecutive strides, it was observed that stride-to-stride fluctuations were not totally  
58 random but instead exhibited a fractal pattern (Hausdorff et al., 1995). Fractal fluctuations in  
59 time series produced by living beings have been deemed to be a signature of their complex

60 internal organization and of the feedback loops needed to adapt behaviors to environmental  
61 changes (Goldberger et al., 2002; West, 2013). Accordingly, physiological time series most often  
62 exhibit scaling properties and statistical persistence. Regarding human walking, the complex  
63 fluctuations in stride intervals, stride speeds, and stride lengths exhibit fractal patterns with  
64 inverse power-law memory (Hausdorff et al., 1995; Terrier, Turner & Schutz, 2005); that is, a  
65 change occurring at a given gait cycle can potentially influence another cycle dozens of steps  
66 later.

67 The fractal pattern of gait fluctuations can be disrupted by sensorimotor synchronization. It is  
68 possible for humans to synchronize their stepping with external rhythmic cues, such as walking  
69 in time with a musical rhythm (auditory cueing). In such cases, stride-to-stride fluctuations  
70 become anti-persistent; that is, stride intervals tend to oscillate stochastically around the imposed  
71 pace (Terrier, Turner & Schutz, 2005; Delignières & Torre, 2009; Sejdić et al., 2012; Choi et al.,  
72 2017). In other words, a long stride interval has a higher probability of being followed by a short  
73 stride interval. Similarly, time series of stride speeds are anti-persistent in treadmill walking, in  
74 which a constant speed is imposed by the treadmill belt (Dingwell & Cusumano, 2010). The  
75 fractal pattern of stride speeds can be restored using self-paced treadmills, in which the belt  
76 speed is dynamically controlled by the walking subjects (Choi et al., 2017). In treadmill  
77 experiments, if an additional instruction of gait synchronization is superimposed on the task of  
78 walking at the belt speed, a generalized anti-persistent pattern is then observed (Terrier & Dériaz,  
79 2012; Roerdink et al., 2015; Choi et al., 2017). This phenomenon exists both when  
80 synchronizing stride intervals to a metronome (auditory cueing), and when aligning step lengths  
81 to visual cues projected onto the treadmill belt (visual cueing) (Terrier, 2016).

82 In 2010, Dingwell and Cusumano hypothesized that the emergence of anti-persistence was  
83 linked to the degree of voluntary control dedicated to the gait. They suggested that, during a  
84 normal gait, deviations go uncorrected and can persist across consecutive strides (under-  
85 correction). In contrast, in paced walking, deviations are followed by rapid corrections that lead  
86 to anti-persistence (over-correction). This “tight control” hypothesis has been supported by other  
87 studies (Roerdink et al., 2015; Bohnsack-McLagan, Cusumano & Dingwell, 2016). Earlier this  
88 year, Roerdink et al. further demonstrated that the degree of anti-persistence can be modulated  
89 by constraining the maneuverability range on a treadmill (Roerdink et al., 2019). In short,  
90 characterizing the noise structure of gait variability helps us to better understand gait control;  
91 among other things, it can provide information about whether a gait is highly controlled or more  
92 automated. In addition, cued walking has important applications for rehabilitation in gait  
93 disorders (Yoo & Kim, 2016; Pereira et al., 2019).

94 Detrended fluctuation analysis (DFA) is typically the preferred method to identify the  
95 fluctuation structure in a time series of gait parameters. Introduced in 1995 by Hausdorff et al.,  
96 DFA identifies the modification of a signal’s variance at different time scales. DFA can unmask  
97 underlying fluctuation structures that may be otherwise obscured by time series trends (Peng et  
98 al., 1995). The presence of power-law scaling is determined through the scaling exponent alpha  
99 ( $\alpha$ ); if the exponent is small ( $\alpha < 0.5$ ), the fluctuations are deemed to be anti-persistent. Statistical

100 persistence corresponds to  $\alpha$  values higher than 0.5 and an  $\alpha$  value equal to 0.5 indicates a  
101 random, uncorrelated noise (see Appendix B in Terrier & Dériaz [2013] for further information).

102 DFA requires a non-periodical, discrete time series as an input. Foot switches, i.e., pressure  
103 sensitive insoles, can be used to detect heel strikes on the ground and can thus collect time series  
104 of stride intervals (Hausdorff, Ladin & Wei, 1995; Sejdić et al., 2012; Almurad et al., 2018).  
105 Several methods using the continuous measure of the positions of various body parts have also  
106 been proposed: 1) high-accuracy GPS (Terrier, Turner & Schutz, 2005); 2) 3-D video analysis  
107 (Dingwell & Cusumano, 2010); and 3) an instrumented treadmill that records the center-of-  
108 pressure trajectory (Terrier & Dériaz, 2012; Terrier, 2016; Roerdink et al., 2019). These methods  
109 require a preliminary discretization of the position signals via minima/maxima detection  
110 algorithms (Terrier & Schutz, 2005; Roerdink et al., 2008; Dingwell & Cusumano, 2010). Other  
111 studies attempted to retrieve stride intervals from acceleration signals (Terrier & Dériaz, 2011),  
112 but the correct discrimination of strides can be challenging (González et al., 2010; Riva et al.,  
113 2013; Terrier & Reynard, 2018).

114 The discrete gait time series that are analyzed through DFA are fundamentally the output of a  
115 continuous process. Indeed, gait control coordinates muscles and joints continuously during  
116 successive gait cycles; this process generates stride intervals, stride lengths, and stride speeds as  
117 outputs. It is questionable whether it is even possible to retrieve the fractal signature of long-  
118 range stride fluctuations in a continuous signal that could capture both intra- and inter- stride gait  
119 dynamics. In 2013, I hypothesized that an attractor that reflects short-term gait dynamics could  
120 also contain information about long-term gait complexity (Terrier & Dériaz, 2013). In 2018, I  
121 explored this hypothesis further (Terrier & Reynard, 2018): I proposed the use of a new gait  
122 complexity index computed from continuous signals, which I named the attractor complexity  
123 index (ACI).

124 ACI is a new term for long-term local dynamic stability (LDS)—also referred to as  
125 divergence exponent or lambda ( $\lambda$ )—which was introduced by Dingwell et al. in 2000 (Dingwell  
126 et al., 2000; Dingwell & Cusumano, 2000). This algorithm, based on Lyapunov exponents used  
127 in chaos theory (Dingwell, 2006; Mochizuki & Aliberti, 2017), has been recommended to assess  
128 gait stability and fall risk (Bruijn et al., 2013). LDS requires the construction of an attractor in  
129 the phase space by means of time delay embedding of continuous signals, such as body  
130 accelerations (Takens, 1981; Rosenstein, Collins & De Luca, 1993; Terrier & Dériaz, 2013).  
131 LDS is defined as the divergence rate among attractor trajectories. The divergence rate can be  
132 evaluated at different intervals, either immediately after the initial separation between adjacent  
133 trajectories (short-term LDS) or several strides later (long-term LDS). In the years following  
134 Dingwell's seminal articles, it became clear that long-term LDS was in fact not a good index for  
135 predicting fall risk and gait stability (Bruijn et al., 2013), but that short-term LDS had better  
136 properties for gait stability analysis, as shown in modeling studies (Su & Dingwell, 2007; Bruijn  
137 et al., 2012).

138 Given that long-term LDS is not a gait stability index, renaming it as ACI seems appropriate.  
139 Indeed, as demonstrated through a modelling approach, ACI is highly sensitive to the noise



140 structure of stride intervals (Terrier & Reynard, 2018). More precisely, a low ACI is associated  
141 with statistical anti-persistence, and a high ACI is associated with persistence. Furthermore, it  
142 has been shown that when stride intervals are kept constant, divergence curves become flat after  
143 only two strides (see Fig. 2 in Terrier & Reynard [2018]). Although additional theoretical work  
144 is required to explore the causes of this sensitivity, it can be assumed that the complex gait  
145 dynamic is reflected by wider boundaries in the attractor, which allows further long-term  
146 divergence. In contrast, statistical anti-persistence signals a less complex gait dynamic, a more  
147 restricted attractor, and therefore a lower long-term divergence rate. The fact that no divergence  
148 is observed if stride intervals are kept constant further supports this hypothesis.

149 The objective of the present study was to confirm that ACI can be used to assess gait  
150 complexity from continuous signals without preliminary discretization. In my 2018 study  
151 (Terrier & Reynard, 2018), I hybridized acceleration signals with artificial signals to explore this  
152 assumption. Here, in order to apply ACI to real signals, I computed both ACI and scaling  
153 exponents ( $\alpha$ s) from a center-of-pressure trajectory recorded in a treadmill experiment that  
154 submitted participants to either visual or auditory cueing. I then explored the responsiveness of  
155 ACI to the cueing conditions, as well as correlations between ACI and  $\alpha$ s. The ability of ACI and  
156  $\alpha$ s to predict cueing conditions was also assessed. The study also had two secondary objectives:  
157 to test the appropriateness of different intervals for computing ACI, and to evaluate short-term  
158 LDS's sensitivity to cueing.

159

## 160 **Materials & Methods**

### 161 **Data**

162 Data from a previous study were re-analyzed (Terrier, 2016). In summary, 36 individuals walked  
163 for 30 min on an instrumented treadmill at their preferred speed. They were exposed to three  
164 different conditions of 10 min duration each: 1) normal walking with no cueing (NC); 2) walking  
165 while synchronizing their gait cadence to an isochronous metronome (auditory cueing, AC); and  
166 3) walking while targeting visually projected shapes with their feet (visual cueing, VC).

### 167 **Ethics statement**

168 The present study is a re-analysis of an anonymized database and is not considered as a human  
169 research needing authorization from an ethic committee. Consent was obtained for  
170 anonymization and reuse. Please refer to the ethic statement in the original publication for further  
171 information (Terrier, 2016).

### 172 **Data availability**

173 Individual data are available in a supplementary file.

### 174 **Data processing**

175 For each condition, 1,000 steps (500 gait cycles) were recorded. The force platform embedded  
176 into the treadmill recorded the position (Cartesian coordinates, anteroposterior [AP] and  
177 mediolateral [ML] axes) of the center of pressure at a sampling rate of 500Hz. Based on the  
178 detection of heel strikes in the anteroposterior (AP) signal, time series of stride time (ST), stride  
179 length (SL) and stride speed (SS) were computed (Roerdink et al., 2008). Next, the noise



180 structure of stride-to-stride fluctuations were assessed with DFA (for in-depth descriptions of the  
181 DFA algorithm, see Terrier, Turner & Schutz [2005] and Terrier & Dériaz [2012, 2013]; DFA  
182 results—the scaling exponents  $\alpha$ —are shown in Terrier [2016]).

183 The 500Hz signal from the AP and ML signals were then low-pass filtered (18Hz 12th order  
184 Butterworth) and down-sampled to 100Hz. After the selection of 300 strides (from the 100th to  
185 the 400th strides), truncated signals were resampled at a constant number of 30,000 samples, i.e.,  
186 100 points per stride.

187 Computations of nonlinear indexes of gait stability (LDS) and complexity (ACI) were  
188 implemented via the same methods as in previous studies that used Rosenstein's algorithm  
189 (Terrier & Dériaz, 2013; Terrier & Reynard, 2015). High dimensional attractors were built  
190 according to the delay-embedding theorem. The average mutual information of each signal was  
191 used to assess the time delay. A common dimension of five was determined with a global false  
192 nearest neighbor analysis. Average divergence of the attractor was defined as  $avg(\ln[d_j(i)])$ , that  
193 is, the logarithm of the  $i^{th}$  Euclidian distance  $d$  downstream of the  $j^{th}$  pair of nearest neighbors in  
194 the attractor, averaged over all pairs. Time was normalized by ST. Resulting divergence curves  
195 are shown in Fig. 1. The exponential divergence rate, calculated as  $avg(\ln[d_j(i)]) / stride$ , was  
196 evaluated with linear fits across several spans as follows: 0–0.5 stride (LDS), 1–4 strides (ACI 1-  
197 4), 4–7 strides (ACI 4-7), and 7–10 strides (ACI 7-10).

#### 198 **Statistics**

199 Notched boxplots were used to depict the distribution of the individual results (Figs. 2 and 3).  
200 Descriptive statistics (means and standard deviations [SD]) were computed for the ACIs (Table  
201 1). LDS statistics can be found in the supplementary file. Fig. 4 shows the effect sizes (Hedges'  
202  $g$ ) of the differences between conditions (i.e., AC minus NC, and VC minus NC), as well as  
203 Bonferroni corrected 95 % confidence intervals.

204 The correlations among the variables are illustrated in Fig. 5 through scatter plots and linear  
205 fits. Pearson's correlation coefficients ( $r$ ) and associated  $p$ -values (null hypothesis for a null  
206 correlation coefficient) were also assessed.

207 Least absolute shrinkage and selection operator LASSO (Tibshirani, 1996) and logistic  
208 regressions were used to assess the extents to which DFA, LDS and ACI could differentiate  
209 between the cueing (AC and VC) and NC conditions. The LASSO algorithm had the advantage  
210 of regularizing the fit for lower overfitting and of selecting the most important predictors. The  
211 dependent binary variable was coded as NC = 1 (36 observations), and AC and VC = 0 (72  
212 observations). Three models were fitted as follows: Model 1: the independent variables were  
213 LDS-AP and LDS-ML (2 predictors); Model 2: the independent variables were ACI 1-4, ACI 4-  
214 7, and ACI 7-10 for both the ML and AP directions (6 predictors); and Model 3: the independent  
215 variables were  $\alpha$ -ST,  $\alpha$ -SL, and  $\alpha$ -SS (3 predictors). All  $\alpha$  values were taken from Terrier (2016).  
216 The LASSO regularization factor was set via 10x cross-validation. Receiver operating  
217 characteristic (ROC) curves were used to illustrate the models' diagnostic abilities. Areas under  
218 the curves (AUCs), along with bootstrapped confidence intervals, were computed as well (Fig.  
219 6). Sensitivity and specificity at  $p = 0.5$  were also evaluated. Fig. 7 presents the standardized

220 coefficients for the three logistic models, which indicate the relative importance of each  
221 predictor.

222

## 223 Results

224 Divergence curves (Fig. 1) revealed a clear difference between cueing and NC conditions,  
225 especially for the AP signal. In the NC condition (black curve), divergence increased steadily,  
226 with moderate dampening. In contrast, for both AC and VC, dampening occurred more rapidly  
227 after four strides.

228 LDS and ACI are defined as slopes of the divergence curves measured at different intervals.  
229 Given the dampening, it was expected that ACI measured further from the initial separation  
230 would exhibit lower values. This was confirmed, as shown in the Fig. 3 boxplots: ACI 1-4 was  
231 higher and more variable than either ACI 4-7 or ACI 7-10. Furthermore, the LDS, which was  
232 computed during the first step, was larger (Fig. 2).

233 As shown by the effect size plots in Fig. 4, ACIs decreased strongly when individuals  
234 followed auditory or visual cues. The effect was most pronounced for the AP signal, for which  
235 both AC and VC had comparable effects. In contrast, a relevant difference existed between NC  
236 and VC for the ML signal.

237 Fig. 5 shows the correlations among the LDS, ACI, and scaling exponents. Of particular note  
238 is the high correlation found between ACI 4-7 measured by the ML direction and the scaling  
239 exponents ( $r = 0.78$  with  $\alpha$ -ST, and  $r = 0.72$  with  $\alpha$ -SL). Other ACI spans exhibited weaker  
240 correlations. ML-LDS was not correlated with other variables, while AP-LDS was weakly  
241 correlated with scaling exponents ( $r = 0.37$  with  $\alpha$ -ST, and  $r = 0.29$  with  $\alpha$ -SL).

242 Using the ACIs and scaling exponents, multivariable logistic models differentiated very well  
243 between the cueing and NC conditions. The AUCs were close to 1 ( $\alpha$  AUC = 0.996, ACI  
244 AUC=0.980; Fig. 6). ACI model's sensitivity was 92% and specificity was 86%. LDS was a  
245 rather poorer predictor (AUC = 0.82, sensitivity = 93%, specificity = 50%).

246 As shown in Fig. 7, The LASSO algorithm selected the most significant predictors, and no  
247 important ones were set to 0. The strongest predictors were  $\alpha$ -ST and ACIs measured in the AP  
248 direction over long-term spans (4-10).

249

## 250 Discussion

251 The aim of this study was to further explore whether ACI could be used to assess gait complexity  
252 from continuous signals. The results strongly support the hypothesis that both DFA and ACI  
253 measure the same thing: their values were strongly correlated, they both differed strongly  
254 between the cueing and NC conditions, and they both predicted cueing conditions with high  
255 degrees of sensitivity and specificity. The results also show that ACI should be measured in the  
256 AP direction and between four to seven strides downstream from the initial separation. In  
257 addition, LDS measured in the ML direction seemed insensitive to cueing, further supporting its  
258 use as a pure gait stability index.

259 A previous study assessed the effect of AC on stride-to-stride fluctuations in a treadmill  
260 experiment among 20 young adults (Terrier & Dériaz, 2012). Scaling exponents of SL and ST  
261 were strongly anti-persistent ( $\alpha < 0.5$ ) under the AC condition. Based on the same data, another  
262 study investigated the effects of AC on LDS and ACI (Terrier & Dériaz, 2013). ACI (still  
263 referred to as  $\lambda$ -L at that time) was computed over a timescale between the 4<sup>th</sup> and 10<sup>th</sup> strides.  
264 The standardized effect size of the difference between the NC and AC conditions was -3.3 for  
265 both the AP and ML signals. In addition, a substantial correlation between the scaling exponents  
266 and ACI was found (canonical correlation:  $r = 0.83$ ). Another research group also found similar  
267 results in a study that combined a foot-switch and an accelerometer to evaluate overground  
268 walking (Sejdić et al., 2012); they found that both ACIs ( $\lambda$ -LT) and scaling exponents were  
269 substantially lower when the walk was paced with a metronome. The results of the present study  
270 confirm ACI's sensitivity to an AC (effect size  $< -2$ ; Fig. 4). Overall, ACI seems sensitive to  
271 changes of long-range fluctuation patterns induced by auditory sensorimotor synchronization.

272 The influence of VC on ACI had not been previously studied. The present results indicate that  
273 both VC and AC induced similar modifications to ACIs measured from the AP signal (Figs. 1  
274 and 4). Previous research has also demonstrated that VC and AC have similar effects on scaling  
275 exponents (Terrier, 2016), which are incidentally computed from the discretization of the AP  
276 signal. In contrast, the present study found that when using ML measures, VC had less of an  
277 effect than did AC (Fig. 4). It is worth noting that the VC procedure consisted of participants  
278 aiming their feet toward rectangular visual targets (stepping stones). As a result, the task required  
279 voluntary leg control in both the AP and ML directions. Further analyses are needed to  
280 specifically explore gait lateral control under such circumstances, for instance by analyzing time  
281 series of step widths, which would be computed from the discretization of the ML signal (see  
282 Terrier, 2012).

283 LDS and ACI are rates of divergence (i.e., slopes) computed from an average logarithmic  
284 divergence curve (Fig. 1). Contrary to a real chaotic attractor, gait divergence curves do not  
285 exhibit a linear region, from which the slope should be computed according to the Rosenstein  
286 algorithm (Rosenstein, Collins & De Luca, 1993; Terrier & Dériaz, 2013). In fact, as illustrated  
287 in Fig. 1, the divergence rate diminishes continuously along the curve. The determination of  
288 range for computing ACI is therefore not straightforward. In their seminal researches, Dingwell  
289 et al. computed the slope between the 4<sup>th</sup> and 10<sup>th</sup> strides, but without a clear justification for this  
290 range (Dingwell et al., 2000; Dingwell & Cusumano, 2000). Subsequent studies followed  
291 identical spans. However, based on an examination of the divergence curves, it may be  
292 unnecessary to go that far from initial separation to estimate a meaningful long-term divergence,  
293 especially since this also increases computational cost. For instance, it was recently shown that  
294 an ACI (LDS-L) computed between the 2<sup>nd</sup> and 6<sup>th</sup> strides could discriminate between healthy  
295 individuals and patients suffering chronic pain of lower limbs (Terrier et al., 2017). In addition,  
296 the recent modeling study that introduced ACI observed that the ACI measured between the 2<sup>nd</sup>  
297 and 4<sup>th</sup> strides was more responsive to the stride-to-stride noise structure than the ACI measured  
298 between the 4<sup>th</sup> and the 10<sup>th</sup> strides, i.e., the originally proposed range (Terrier & Reynard, 2018).

299 Here, the results show that ACI 4-7 was superior to the other ranges: it exhibited the highest  
300 correlation with the scaling exponents of ST and SL ( $r = 0.78$  and  $0.72$ ; Fig. 5), it had the highest  
301 contrast with the NC condition (Fig. 4), and it was selected by the logistic model as the second  
302 highest predictor of cueing conditions (standardized coefficient =  $0.89$ ; Fig. 7). In short, it is very  
303 likely that it is not necessary to measure divergence after the 7<sup>th</sup> stride to assess ACI.

304 The results also indicate that LDS did not respond similarly in the ML and AP directions.  
305 Indeed, ML-LDS was not correlated with complexity measures (ACI and  $\alpha$ ) and had no  
306 predictive power (Fig. 7). In contrast, AP-LDS was moderately, but significantly, correlated with  
307 complexity measures ( $r = .37$  and  $.29$ , Fig. 5) and was solely responsible for the LDS model's  
308 moderate capacity to differentiate between cueing conditions (AUC =  $.82$ ; Fig. 7 and 8). ML-  
309 LDS has been shown to be an index of gait instability (Reynard et al., 2014) and fall risk  
310 (Bizovska et al., 2018). This may be due to the importance of lateral stability for maintaining a  
311 steady and safe gait (Bauby & Kuo, 2000; Gafner et al., 2017). The results of the present study  
312 support the use of ML-LDS for stability assessments given its independence from complexity  
313 measures. In contrast, it can be assumed that interactions exist between the long-term noise  
314 structure of a gait and its short-term stability in the AP direction; this lack of independence may  
315 obscure the significance of the AP-LDS measure. However, it is unclear whether results obtained  
316 from center-of-pressure trajectory are comparable to those obtained with other methods, such as  
317 trunk accelerometry; incidentally, a large-scale accelerometry study found that AP-LDS could  
318 predict future falls (van Schooten et al., 2015). The assumption that ML-LDS is better suited for  
319 gait stability assessments thus requires further investigations.

320 The biggest strength of the present study is in its substantial number of strides measured in a  
321 large sample of healthy adults (36), particularly when compared to other recent studies in the  
322 field (Sejdić et al., 2012; Bohnsack-McLagan, Cusumano & Dingwell, 2016; Roerdink et al.,  
323 2019). Evaluating gait complexity requires the analysis a large number of consecutive strides  
324 (Marmelat & Meidinger, 2019). Similarly, reliability results show that many consecutive strides  
325 are required to accurately assess ACI (Reynard & Terrier, 2014). Consequently, this study's  
326 findings most likely offer good generalizability. The study's primary limitation is that the  
327 analyses of the center-of-pressure trajectories are restricted to treadmill experiments with few  
328 potential applications. The center-of-pressure approach has the advantage of allowing an easy  
329 discretization to compare both discrete time series and continuous signals (Roerdink et al., 2008),  
330 but further investigations are required to explore ACI potential in real-life applications using  
331 inertial sensors such as accelerometers.

332

## 333 **Conclusions**

334 This study's findings support the hypothesis that ACI can provide information about the stride-  
335 to-stride fluctuation structure of an individual's gait based on continuous signals. Accordingly,  
336 information about gait complexity can be obtained while measuring a gait with inertial sensors,  
337 such as accelerometers (Terrier et al., 2017; Terrier & Reynard, 2018). ACI could thus assess the  
338 degree of motor control applied by walkers on their gait (the "thigh control" hypothesis; see

339 Dingwell & Cusumano [2010] and Roerdink et al. [2019]). A high ACI would indicate an  
340 automated gait, while a lower ACI would be a sign of greater voluntary attention dedicated to  
341 gait control. For example, it has been previously suggested that a low ACI in patients with lower  
342 limb pain is due to enhanced gait control to avoid putting too much weight on a painful leg  
343 (Terrier et al., 2017). Older studies that inappropriately used ACI as a gait stability index should  
344 be reinterpreted with the “thigh control hypothesis” taken into account. For example, Dingwell  
345 et al. found that patients suffering from peripheral neuropathy had lower ACIs, which was  
346 interpreted as a higher gait stability obtained by lowering walking speed (Dingwell et al., 2000).  
347 An alternative explanation would be that diminished sensory feedback required more attention  
348 dedicated to gait control.

349 The use of LDS to characterize gait stability and assess fall risk has gained popularity over  
350 recent years (Mochizuki & Aliberti, 2017; Bizovska et al., 2018; Mehdizadeh, 2018). Computing  
351 ACI in addition to LDS is straightforward and using the measures together could be fruitful, as  
352 information about gait automaticity and cautiousness would complement information about gait  
353 stability. It is hoped that the results of this study will help convince future researchers to reinstate  
354 the use of ACI to further enrich their gait analysis studies.

355

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357 None

358

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**Table 1** (on next page)

Table 1: Descriptive statistics of the attractor complexity index (ACI)

Means and standard deviations (SD) of ACI measured in the 36 subjects under the three experimental conditions. AP: anteroposterior. ML: mediolateral.

1 **Table 1: Descriptive statistics of attractor complexity index (ACI)**

2

ACI x 1000	<b>N=36</b>		<b>ACI 1-4</b>				<b>ACI 4-7</b>				<b>ACI 7-10</b>			
	AP		ML		AP		ML		AP		ML			
	mean	SD	mean	SD	mean	SD	mean	SD	mean	SD	mean	SD		
No cueing	2.00	(0.31)	1.25	(0.33)	0.78	(0.12)	0.48	(0.17)	0.44	(0.13)	0.34	(0.12)		
Auditory cueing	1.55	(0.36)	0.89	(0.21)	0.42	(0.17)	0.17	(0.12)	0.18	(0.16)	0.07	(0.11)		
Visual cueing	1.29	(0.43)	1.01	(0.33)	0.34	(0.23)	0.31	(0.20)	0.14	(0.14)	0.15	(0.15)		

3

4 Means and standard deviations (SD) of ACI measured in the 36 subjects under the three  
5 experimental conditions. AP: anteroposterior. ML: mediolateral.

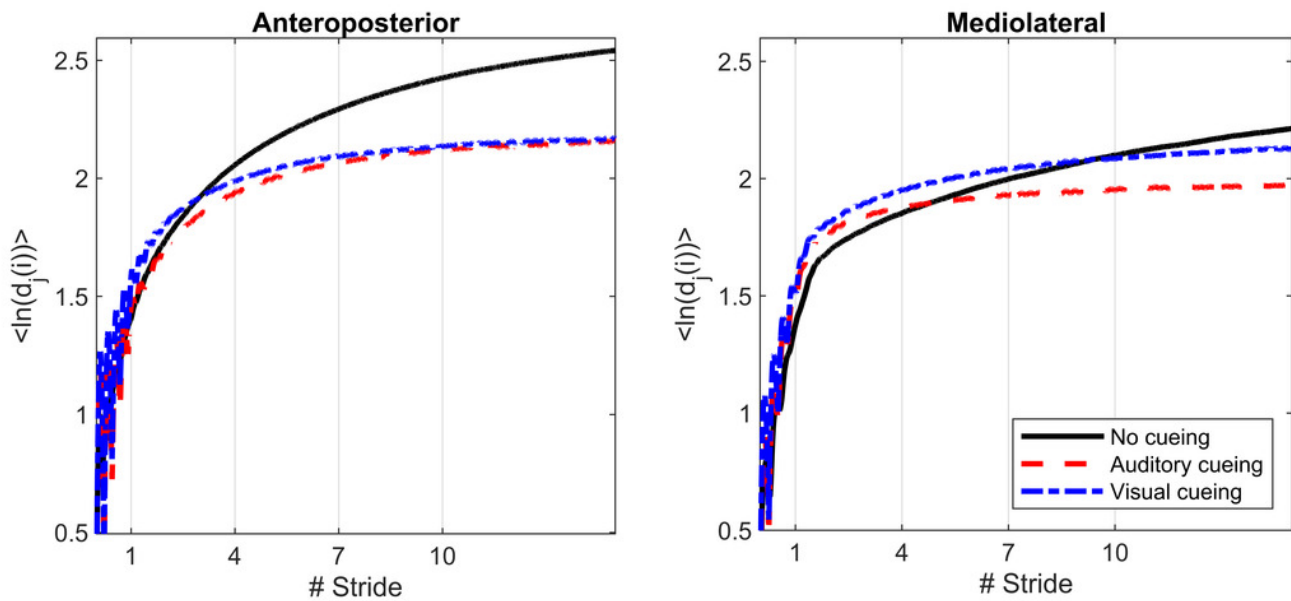
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7

# Figure 1

Figure 1: Divergence curves

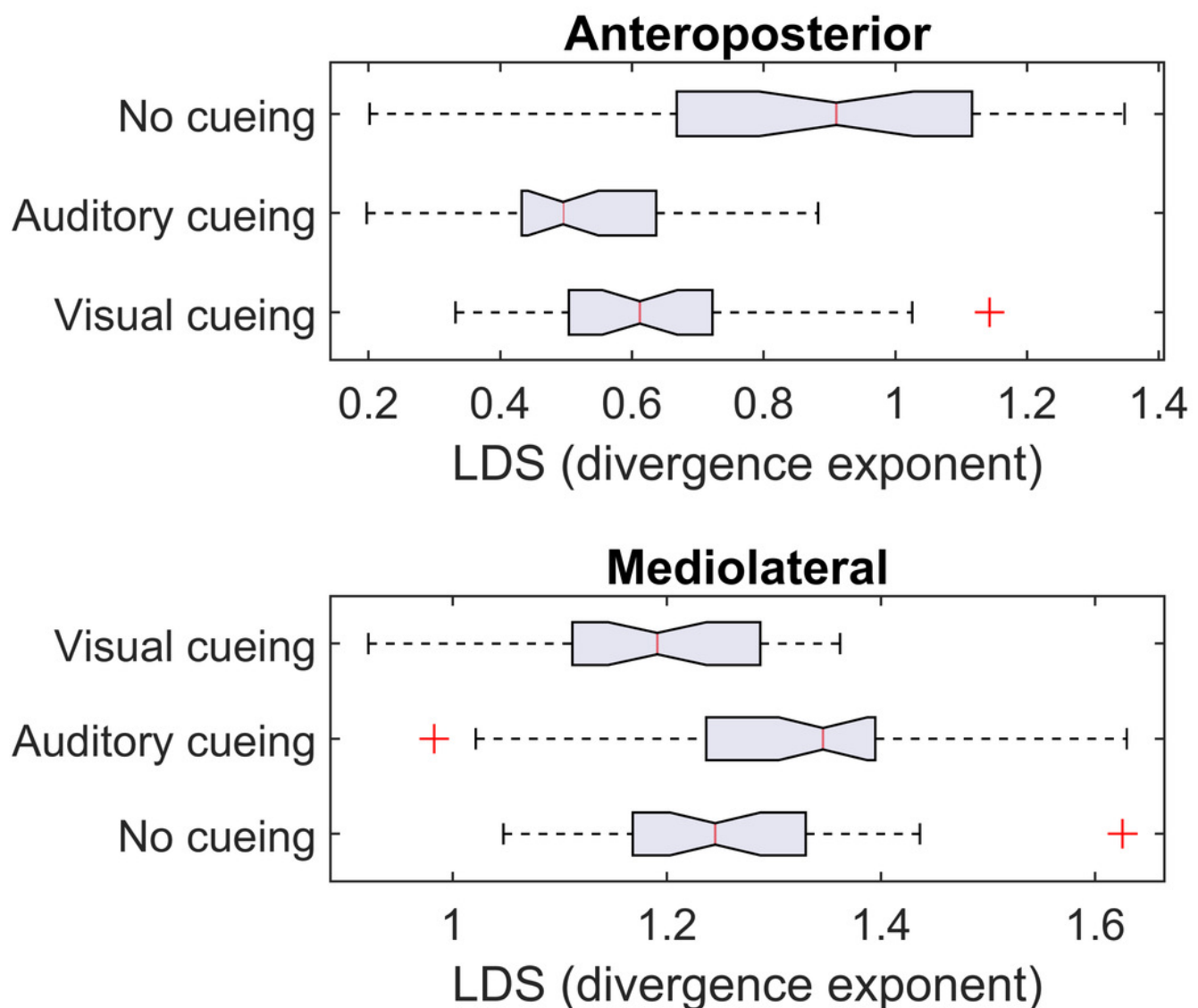
Using time-delay embedding, 5-dimensional attractors were reconstructed from the anteroposterior and mediolateral coordinates of a center-of-pressure trajectory. The logarithmic divergence from neighbor trajectories (y-axis) was averaged across trajectories and participants (N=36), and drawn against normalized time (strides, x-axis). Three curves are shown, one for each experimental condition.



## Figure 2

Figure 2: Descriptive statistics of the local dynamic stability (LDS)

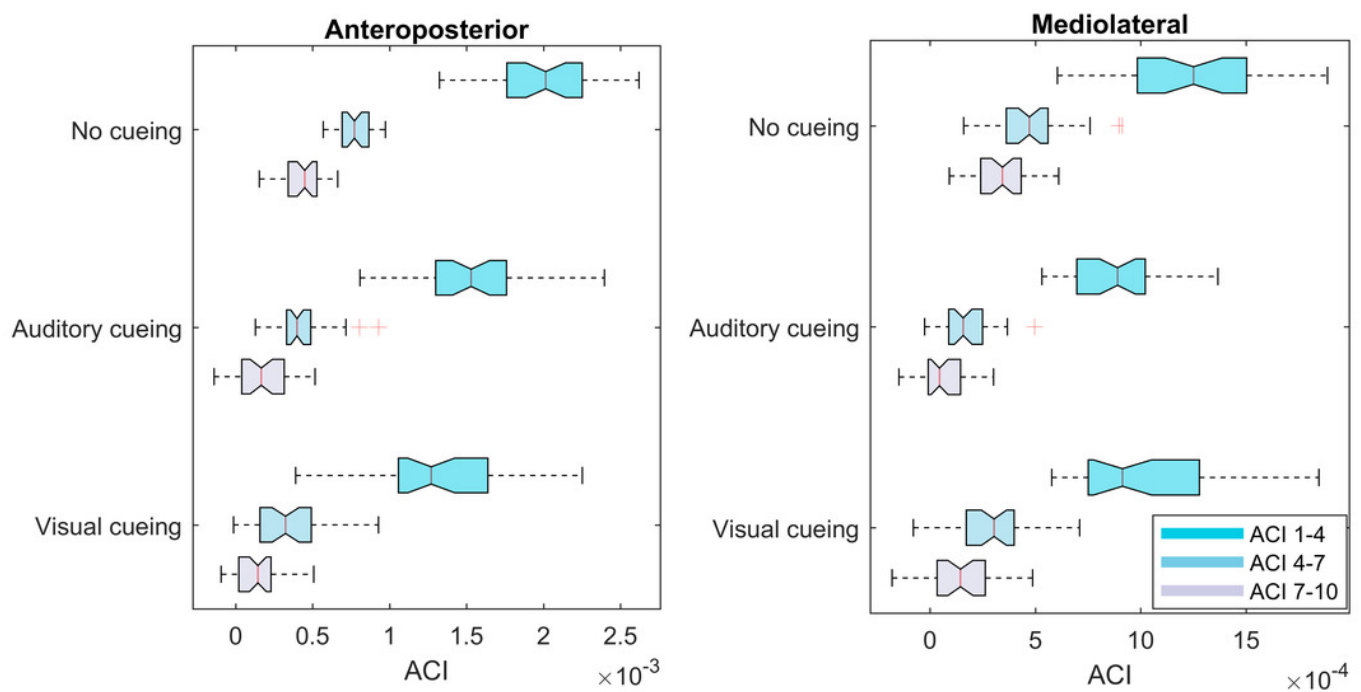
The notched boxplots summarize the distribution of individual results ( $N = 36$ ) across the three experimental conditions. The notch extremes correspond to the 95% confidence intervals of the medians. The red + symbols indicate outliers.



## Figure 3

Figure 3: Descriptive statistics of the attractor complexity index (ACI)

The notched boxplots summarize the distribution of individual results ( $N = 36$ ) across the three experimental conditions for the three different ACI spans. The notch extremes correspond to the 95% confidence intervals of the medians. The red + symbols indicate outliers.

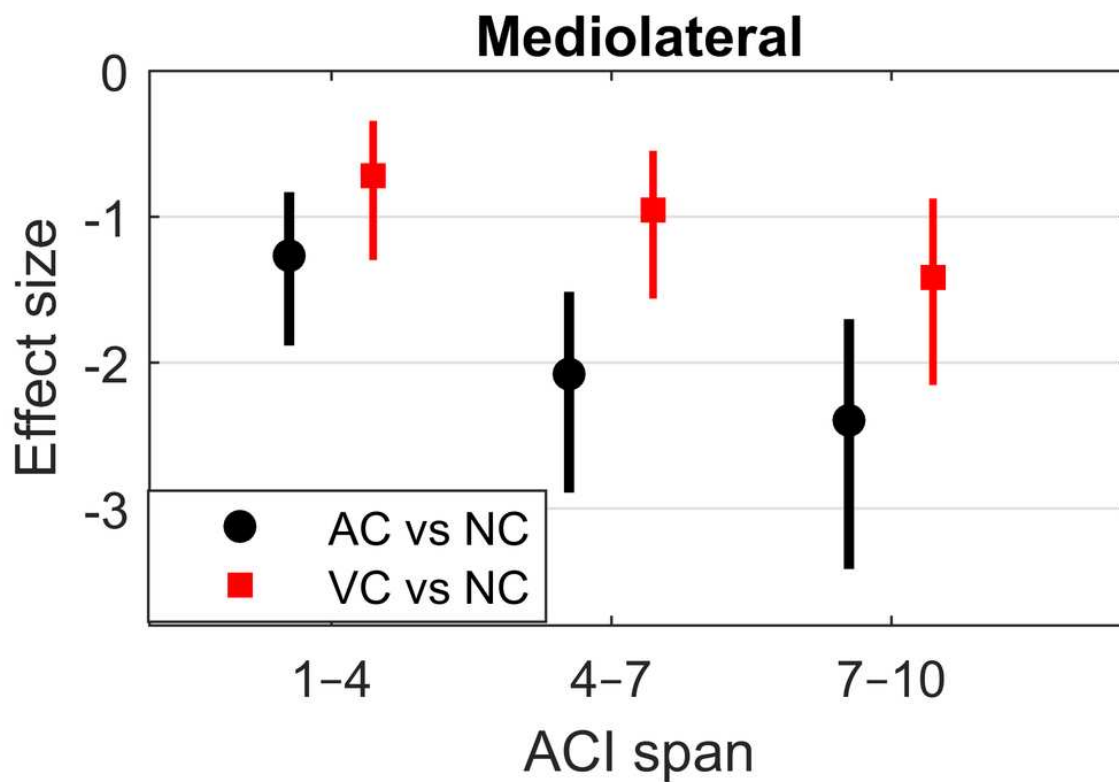
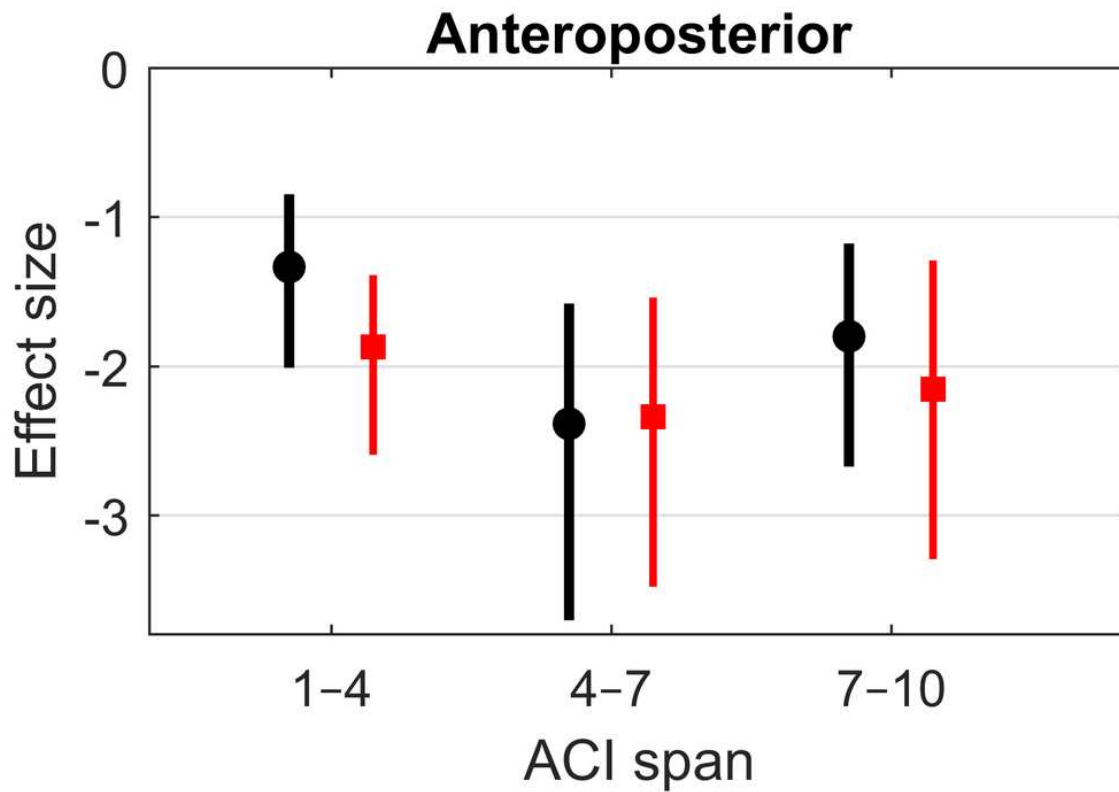


## Figure 4

Figure 4: Effect sizes of attractor complexity index (ACI)

Standardized effect size (Hedges'  $g$ ) of the difference between cueing and no-cueing conditions. Vertical lines are 95% confidence intervals (Bonferroni corrected). AC: auditory cueing; VC: visual cueing; NC: no-cueing.

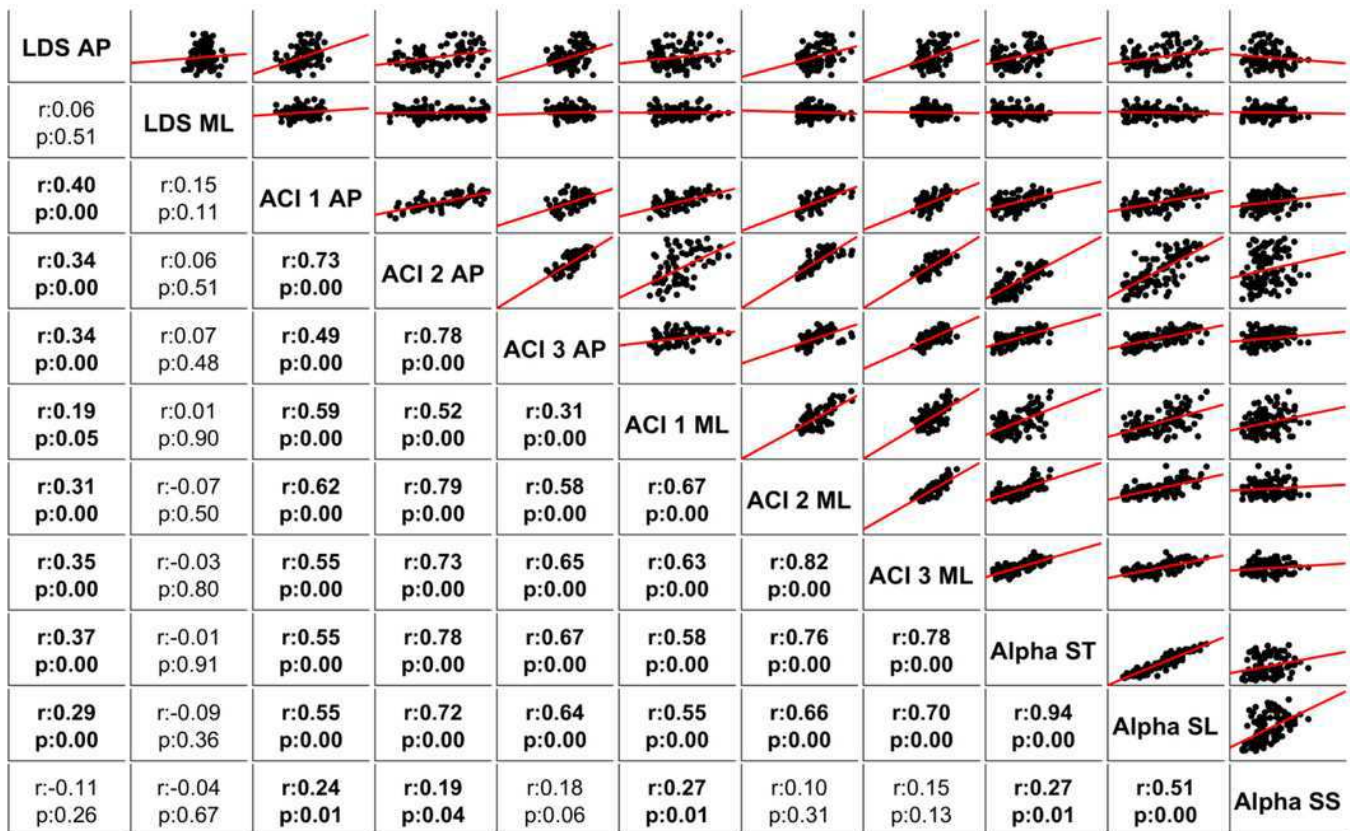




## Figure 5

Figure 5: Correlations and scatter plots across local dynamic stability (LDS), attractor complexity index (ACI), and scaling exponent (alpha) measures

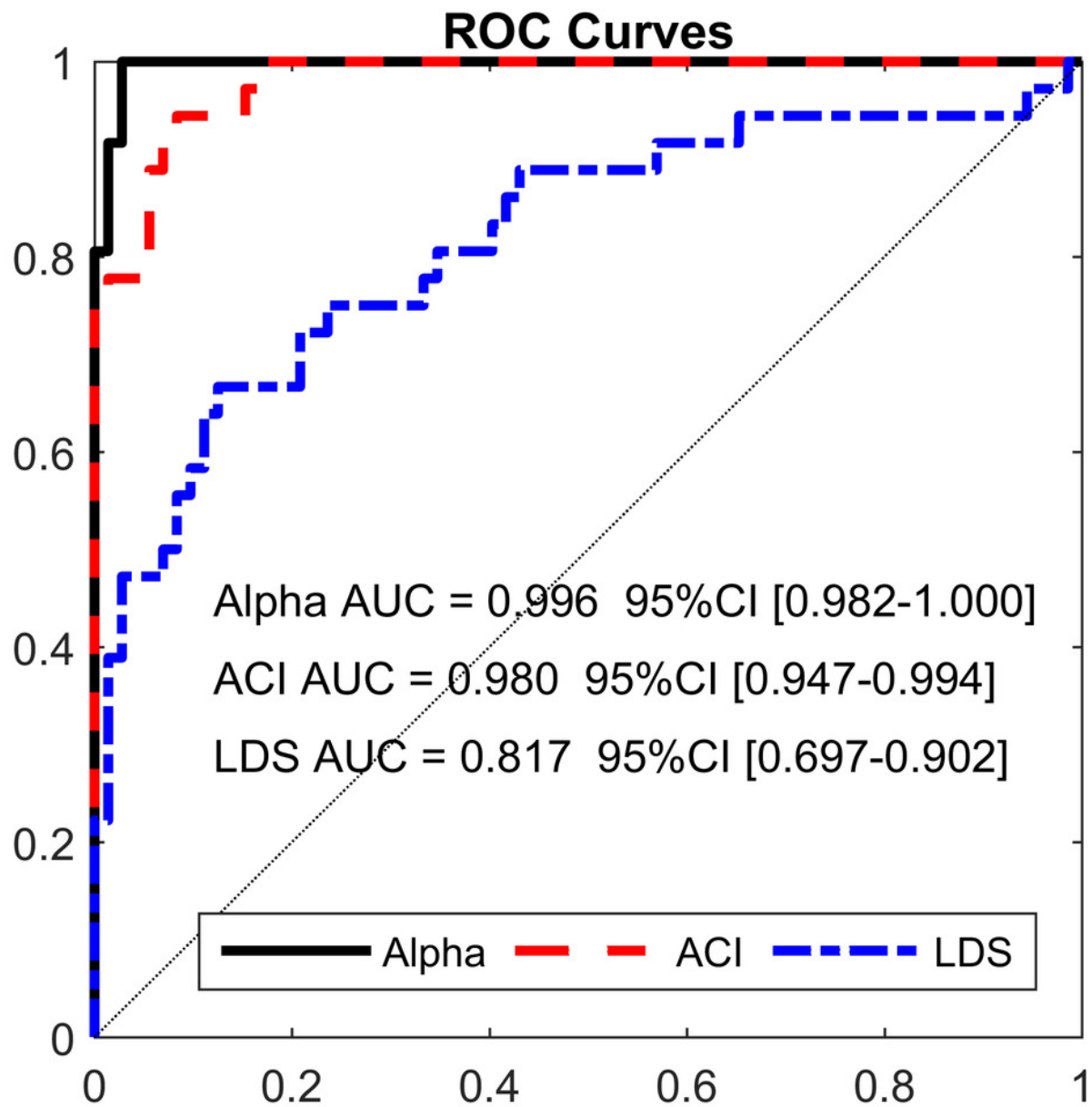
Pearson's correlation coefficients ( $r$ ) are shown on the lower left, along with the results for the hypothesis test for  $r = 0$ . Bold values indicate significant results. In the upper right, scatter plots with the linear fits are shown. AP: anteroposterior; ML: mediolateral; ST: stride time; SL: stride length; SS: stride speed.



## Figure 6

Figure 6: Receiver operating characteristic (ROC) curves

ROC curves for the three multivariable logistic models predicting cueing/no-cueing conditions: 1) local dynamic stability (LDS); 2) attractor complexity index (ACI); and 3) scaling exponent (alpha). Areas under the curves (AUCs) are written with their confidence intervals.



## Figure 7

Figure 7: Standardized coefficients of the multivariable logistic models

Three multivariable logistic models were fitted: 1) local dynamic stability (LDS); 2) attractor complexity index (ACI); and 3) scaling exponent (Alpha). A least absolute shrinkage and selection operator (LASSO) was used to regularize fitting. Bars show the value of the standardized beta coefficient of the regressions for each predictor. AP: anteroposterior; ML: mediolateral; ST: stride time; SL: stride length; SS: stride speed.

