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# Introducing Statistical Persistence Decay – A Quantification of Stride-to-Stride Time Interval Dependency in Human Gait

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1 **ORIGINAL RESEARCH ARTICLE**

2  
3 **Title:**

4 Introducing Statistical Persistence Decay – A Quantification of Stride-to-Stride Time Interval  
5 Dependency in Human Gait  
6

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1 **Abstract**

2 Stride-to-stride time intervals during human walking is characterised by predictability and  
3 statistical persistence quantified by sample entropy (SaEn) and detrended fluctuation analysis  
4 (DFA) which indicates a time dependency in the gait pattern. However, neither analyses quantify  
5 time dependency in a physical or physiological interpretable time scale. Recently, entropic half-  
6 life ( $ENT^{1/2}$ ) has been introduced as a measure of the time dependency on an interpretable time  
7 scale. A novel measure of time dependency, based on DFA, statistical persistence decay (SPD),  
8 was introduced. The present study applied SaEn, DFA,  $ENT^{1/2}$ , and SPD in known theoretical  
9 signals (periodic, chaotic, and random) and stride-to-stride time intervals during overground and  
10 treadmill walking in healthy subjects. The analyses confirmed known properties of the  
11 theoretical signals. There was a significant lower predictability ( $p=0.033$ ) and lower statistical  
12 persistence ( $p=0.012$ ) during treadmill walking compared to overground walking. No significant  
13 difference was observed for  $ENT^{1/2}$  and SPD between walking condition, and they were  
14 exhibited a low correlation.  $ENT^{1/2}$  showed that predictability in stride time intervals was halved  
15 after 11–14 strides and SPD indicated that the statistical persistency was deteriorated to  
16 uncorrelated noise after  $\sim 50$  strides. This indicated a substantial time memory, where  
17 information from previous strides affected the future strides.

18

19

20 **Keywords:** walking, dynamics, nonlinear behaviour, entropy, DFA, stride time fluctuations

21

22

## 1 Introduction

2 The stride time pattern during continuous walking in healthy individuals has been shown to  
3 include stride-to-stride fluctuations exhibiting statistical persistence<sup>6,14</sup>. Thus, each stride  
4 depends on many previous strides with a stride-to-stride dependency that “*decay in a scale-free*  
5 *(fractal-like), power-law fashion*”<sup>12</sup>. Equally, stride length fluctuations during both treadmill and  
6 overground walking have been shown to exhibiting statistical persistence<sup>6,32</sup>. While stride speed  
7 fluctuations during treadmill walking have been shown to exhibit statistical anti-persistence<sup>6</sup>,  
8 statistical persistence has been observed during overground walking<sup>32</sup>.

9 Reduced stride-to-stride persistence has been interpreted differently in relation to the function of  
10 the underlying motor control system. Loss of statistical persistence in stride time fluctuations has  
11 been observed in older adults<sup>13,16</sup>, different neurological patients<sup>11,13</sup>, and has been suggested to  
12 reflect a degraded motor control function<sup>10</sup>. However, reduced persistence observed in frail  
13 individuals has been suggested to potentially implicate an increased control effort to achieve a  
14 more cautious gait pattern<sup>6</sup>.

15 These contradicting observations call for caution when linking loss of stride-to-stride persistence  
16 to either an impaired motor control function or enhanced motor control effort<sup>6</sup>. The  
17 aforementioned studies have applied detrended fluctuation analysis (DFA) to assess the presence  
18 and strength of statistical persistence in the investigated time series. However, DFA does not  
19 quantify the time dependency on an interpretable physiological or physical time scale. This  
20 means that even though existence of statistical persistence can be confirmed by DFA, the stride-  
21 to-stride dependency cannot be quantified in terms of a specific number of previous strides that  
22 influences the current stride. Equally, DFA does not quantify for how long into the future in  
23 terms of seconds or minutes a completed stride will influence new strides.



1 In addition to DFA, sample entropy (SaEn) has been applied to both kinematic signals and stride  
2 time interval time series recorded during gait in order to quantify the predictability of the gait  
3 pattern<sup>1,9,18</sup>. SaEn is high in random white noise signals where no point-to-point dependency  
4 exists and the predictability is low. In contrast, in both chaotic and periodic signals, point-to-  
5 point dependency does exist and they are characterized by relative low SaEn indicating high  
6 predictability (e.g. demonstrated in<sup>33</sup>). However, the predictability is reported on a relative scale  
7 and cannot be translated into physical or physiological terms. Both DFA and SaEn quantify the  
8 time dependency of a time series but return outcome values not easily interpretable in relation to  
9 other physiological measurements (e.g. duration of muscle activity, latency in reflex  
10 measurements, reaction time).

11 Inspired by multiscale entropy (MSE) and to overcome the aforementioned methodological  
12 limitation, entropic half-life ( $ENT^{1/2}$ ) was proposed by Zandiyeh and Von Tscharner<sup>34</sup>.  $ENT^{1/2}$   
13 estimates the time until the predictability in a time series is halved. This is also a measure of how  
14 long data points remain related to one another. Applied to movement related variables,  $ENT^{1/2}$   
15 could quantify how long time elapses before previously performed movements have substantially  
16 reduced their influence on future movements<sup>2</sup>.

17 While DFA, SaEn, and MSE previously have been applied to characterize the stride-to-stride  
18 time dependency in human gait,  $ENT^{1/2}$  has to the best of our knowledge not previously been  
19 used to quantify time dependency in stride-to-stride time intervals. When applied to stride-to-  
20 stride time interval time series,  $ENT^{1/2}$  will estimate how long (in terms of number of strides) it  
21 takes to deteriorate the predictability of the stride time intervals by 50 %. DFA could be used to  
22 quantify decay in time persistence of time series through an application similar in manner to  
23  $ENT^{1/2}$ .

1 In many gait experiments, the treadmill has been used instead of overground walking due to its  
2 advantages with respect to continuous data collection of motion capture, ground reaction forces,  
3 etc. The results and interpretation have often been extrapolated to overground walking even  
4 though substantial biomechanical differences have been observed between treadmill and  
5 overground walking<sup>20,31</sup>. The constraints of the constant speed and limited space of the treadmill  
6 have been suggested to induce a less persistent and more unpredictable walking pattern  
7 compared to overground walking<sup>31</sup>.

8 The present study aimed at introducing two novel tools, ENT<sup>1/2</sup> and SPD as methods to quantify  
9 time dependency in stride-to-stride time intervals during human gait. To validate the use of these  
10 methods, the present study included known theoretical signals (i.e., periodic, chaotic, and  
11 random) and stride time data recorded during overground and treadmill walking. With respect to  
12 the theoretical signals, it was hypothesized that the periodic signals would be characterized by  
13 high ENT<sup>1/2</sup> and SPD, the random signal would be characterized by low ENT<sup>1/2</sup> and SPD and the  
14 chaotic signals would be characterized by intermediate ENT<sup>1/2</sup> and SPD.

15 Further, experimental data from one-hour of walking overground and on a treadmill were used to  
16 verify the use of the methods. Based on previous observations of less statistical persistency and  
17 more unpredictability during treadmill walking<sup>31</sup>, we hypothesized that the time dependency  
18 during treadmill will be less pronounced indicated by lower ENT<sup>1/2</sup> and SPD values compared to  
19 overground walking.

20 As a secondary aim, the association between statistical persistence and predictability were  
21 investigated. If changes in the statistical persistence in stride-to-stride time intervals observed  
22 during human walking<sup>6,14</sup> could explain changes in the predictability; it was hypothesized that  
23 the scaling exponent and SaEn in such time series would correlate. Furthermore, if changes in

1 the deterioration in statistical persistence in stride-to-stride time intervals could explain changes  
2 in the deterioration in the predictability,  $ENT^{1/2}$  and SPD values would also correlate. Such  
3 correlations could indicate that the underlying mechanisms for creating statistical persistence and  
4 predictability are regulated simultaneously.

5

## 1 **Materials and Methods**

### 2 **Theoretical Procedures**

3 To verify the interpretation of the outcome measure, twenty time series of four different  
4 mathematical signals of 2500 data points with different characteristics (periodic, chaotic, and  
5 random) were created using the *colored noise generator* function in Matlab (MathWorks  
6 R2011b). These signals included a brown noise signal (power spectrum of  $1/f^2$ ) which was  
7 considered to be periodic, a signal derived from the second Lorenz differential equation ( $\sigma = 10$ ,  
8  $\rho = 28$ , and  $\beta = 8/3$ ) which was considered to be chaotic, a pink noise signal (power spectrum of  
9  $1/f$ ) which also was considered to be chaotic, and a white Gaussian noise signal (constant power  
10 spectrum) which was considered to be random (figure 1). The periodic brown noise signal and  
11 the random Gaussian noise signal represented two extremes for each of the four applied analyses  
12 with the chaotic Lorenz attractor and pink noise as intermediate signals.

### 13 **Subjects**

14 Fourteen volunteers (seven males and seven females) with a mean ( $\pm$  SD) age of 25.0 years ( $\pm$   
15 4.2), height of 170.8 cm ( $\pm$  11.9) and body mass of 69.4 kg ( $\pm$  16.9) participated in the present  
16 study. The participants had no diagnosed lower limb injuries within the past year. They were  
17 informed of the experimental conditions and gave their written consent to participate in the  
18 study. The study was approved by the by the Institutional Review Board of the University of  
19 Nebraska Medical Center, and it was carried out in accordance with the approved guidelines.

### 20 **Protocol**

21 The study consisted of two experimental sessions. During the first session the subjects completed  
22 a one hour overground walking trial on an elliptical indoor track (circumference  $\sim$  201m) at their  
23 self-selected walking speed. The walking speed was not recorded and was allowed to fluctuate.

1 At the second session, the subjects completed a one hour treadmill walking trial at a constant  
2 self-selected walking speed. The walking speeds were not registered. During both walking trials,  
3 footswitches (Trigno™ 4-channel FSR Sensor, Delsys Inc., Natick, MA) placed under both heels  
4 recorded heel strikes at a sampling rate of 148 Hz. No objective measurement of fatigue was  
5 obtained during the trials. However, none of the subjects reported fatigue to influence their gait.

## 6 **Analysis**

7 The right heel strike data series from each walking trial was processed in Matlab (MathWorks  
8 R2011b) in order to create stride time interval time series (figure 1 and supplementary material  
9 1). Stride time was defined as the time from heel strike of one foot until the subsequent heel  
10 strike of the same foot. Each time series was cut to contain 2500 strides and were subjected to  
11 four different analyses: 1) SaEn, 2) ENT<sup>1/2</sup>, 3) DFA, and 4) SPD.

### 12 *Sample entropy*

13 SaEn was based on the algorithm by Richman and Moorman<sup>27</sup> (equation 1). SaEn was defined as  
14 the negative logarithm for conditional properties that a series of data points within a certain  
15 distance,  $m$ , would be repeated within the distance  $m+1$ <sup>27</sup>.

$$16 \quad \textbf{Equation 1: } SaEn(m, r, N) = -\ln\left[\frac{A^{m+1}(r)}{B^m(r)}\right]$$

17 Where  $A$  is the number of similar vector lengths ( $m+1$ ) falling within a relative tolerance limit ( $r$   
18 times standard deviation of the stride time intervals) and  $B$  is the number of similar vector  
19 lengths ( $m$ ) falling within the tolerance limit<sup>33</sup>. The three parameters  $m$ ,  $r$ , and  $N$  (time series  
20 length) should be selected prior to calculating SaEn and have been shown to have crucial  
21 importance for the SaEn value<sup>33</sup>. In order to control for parameter consistency, SaEn was  
22 calculated using  $m$  of 2 and 3 and  $r$  of 0.05, 0.1, 0.15, 0.2, 0.25, and 0.3. Based on this analysis

1 (see supplementary material 2) m of 2 and r of 0.2 were used for both the SaEn and ENT<sup>1/2</sup>  
2 analyses.

### 3 *Entropic half-life*

4 ENT<sup>1/2</sup> is based on consecutive calculations of SaEn with increasing randomization of the stride  
5 time interval time series as described briefly below and in detail elsewhere<sup>2,34</sup>. Firstly, SaEn is  
6 calculated on the original time series. Secondly, the original time series is gradually randomised  
7 through successive reshaping according to the principle described in figure 2. In the present  
8 study, each reshaping resulted in an increased distance between two subsequent strides. The  
9 stride time interval time series was reshaped 100 times and SaEn was calculated for each of the  
10 reshaped time series. These reshaped time series would, in all cases, exhibit SaEn values  
11 between the SaEn of the original time series (lowest SaEn value) and the SaEn of a complete  
12 random time series (highest SaEn value). The SaEn of the reshaped time series were normalized  
13 to the difference between these two extremes according to equation 2:

14 **Equation 2:** 
$$\text{Normalized SaEn} = \frac{\text{SaEn}_{RS} - \text{SaEn}_{OR}}{\text{SaEn}_{RAN} - \text{SaEn}_{OR}}$$

15 where SaEn<sub>RS</sub> is the SaEn of the reshaped time series, SaEn<sub>OR</sub> is the SaEn of the original time  
16 series, and SaEn<sub>RAN</sub> is the average SaEn of 100 randomized time series created by a random  
17 permutation of the data points in the original time series.

18 The normalized SaEn values were then plotted in a semi logarithmic plot as a function of the  
19 stride number (figure 2). The stride number corresponding to the first SaEn value above 0.5 was  
20 considered the stride number indicating a change in the characteristics of the reshaped time series  
21 from predictability to unpredictability and termed entropic half-life<sup>2,34</sup>.

### 22 *Detrended fluctuation analysis*

1 The presence of statistical persistence or anti-persistence in the stride time interval time series  
2 was assessed using DFA. The correlations related to persistence or anti-persistence are part of  
3 the multifractal cascades that exist over a wide range of time scales<sup>14</sup>. DFA has the advantage of  
4 enabling detection of statistical persistence within noisy signals with embedded polynomial  
5 trends<sup>24</sup>. The applied DFA algorithm has been described in details elsewhere<sup>24</sup> and briefly below.  
6 Using equation 3, the time series  $x(i)$  is first integrated by calculating the cumulated sum of the  
7 deviations of the mean

8 **Equation 3:** 
$$y(k) = \sum_i^k [x(i) - x_{ave}]$$

9 Next, the time series is divided into boxes of equal length,  $n$  and a least square line is fitted to  
10 each box. The  $y$  coordinate of the straight-line segments is designated by  $y_n(k)$  and used to  
11 detrend the time series  $y(k)$  before the root mean square is calculated (equation 4).

12 **Equation 4:** 
$$F(n) = \sqrt{\frac{1}{N} \sum_{k=1}^N [y(k) - y_n(k)]^2}$$

13 This procedure is repeated across the entire time series in order to establish a relationship  
14 between the average fluctuation,  $F(n)$ , as a function of box size  $n$ . The fluctuations can be  
15 characterized by the scaling exponent, which is determined by finding the slope of the line  
16 relating  $\log F(n)$  to  $\log n$ <sup>24</sup>. In the current study, a box size range of  $[2, N]$  and a scaling region of  
17  $10 - 30$  were used for the DFA as this range represented a linear section of the  $\log F(n)$ - $\log n$   
18 plot<sup>14</sup>. A scaling exponent greater than 0.5 indicated a presence of statistical persistence meaning  
19 that a deviation in stride time from the mean in one direction is more likely to be followed by a  
20 deviation in stride time in the same direction. A scaling exponent less than 0.5 indicated a  
21 presence of statistical anti-persistence, meaning that a deviation in stride time from the mean in  
22 one direction is more likely to be followed by a deviation in stride time in the opposite direction.

1 If the scaling exponent is 0.5, this indicated an uncorrelated white noise like pattern of the stride  
2 time interval time series<sup>6,13,14</sup>.

### 3 *Statistical persistence decay*

4 The reshaped time series used for ENT<sup>1/2</sup> analysis were also used for SPD. For each reshaped  
5 time series, DFA was performed as described above with a box size range of [2,N] and a scaling  
6 region of 10 – 30. A critical limit was calculated following equation 5 (figure 3).

7 **Equation 5:**  $Critical\ limit = \mu_{\alpha Ran} + 2 \cdot \sigma_{\alpha Ran}$

8 Where  $\mu_{\alpha Ran}$  is the averaged scaling exponent of 100 random time series created by a random  
9 permutation of the data points in the original time series and  $\sigma_{\alpha Ran}$  is the corresponding standard  
10 deviation. Thus, the critical limit is based on the upper 95% confidence limit ( $\mu + 2\sigma$ ) of the  
11 scaling exponent of the randomised time series. As the order of data points is changed with every  
12 rescaled time series, the statistical persistence is changed towards the critical limit. Any scaling  
13 exponent below the critical limit will not be significantly different from that of randomised  
14 patterns in the original time series. The number of strides required to reduce the scaling exponent  
15 below the critical limit is considered the SPD. Thus, SPD indicates a change in the stride time  
16 interval fluctuation from statistical persistence towards uncorrelated noise.

### 17 **Statistics**

18 Statistical difference in the analysis outcome measures (scaling exponent, SaEn, ENT<sup>1/2</sup>, and  
19 SPD) between the four mathematical signals were assessed using a one way ANOVA on ranks  
20 with signal types as independent factor and the outcome measures as the dependent measure. In  
21 case of an overall significant effect, a Turkey post hoc test was applied.

22 Paired Student's t-tests were applied to SaEn, ENT<sup>1/2</sup>, scaling exponent, and SPD to investigate if  
23 there was a statistical difference between the two walking conditions. To determine the nature of



1 the linear relationship between scaling exponent and SaEn and between  $ENT^{1/2}$  and SPD during  
2 both overground and treadmill walking, linear regression analyses were applied. For all statistical  
3 analysis the level of significance was set at 5%. All statistical calculations were performed in  
4 Sigmaplot (Systat Software, Inc. 2014, version 13.0, Germany).

5

## 1 **Results**

### 2 **Theoretic signals**

3 The scaling exponent, SaEn, ENT<sup>1/2</sup> and SPD of the brown noise, Lorenz attractor, pink noise,  
4 and white Gaussian noise signals are presented in table 1. As known signals, the brown and  
5 Gaussian noise signals represented the two extremes (periodic and random, respectively) with  
6 the chaotic Lorenz attractor and pink noise signals as intermediate signals. The four analyses  
7 confirmed the known properties of the four theoretical signals. For all four analyses, there was a  
8 significant overall effect of the type of signal ( $p < 0.001$  in all cases). The post hoc analyses  
9 revealed a significant difference between each signal type for the scaling exponent ( $p \leq 0.034$  in  
10 all cases). SaEn did not differ between brown noise and the Lorenz attractor but was significantly  
11 higher for pink and Gaussian noise ( $p \leq 0.033$  for all comparisons). ENT<sup>1/2</sup> differed significantly  
12 between all signals ( $p \leq 0.033$  for all comparisons) except between pink and Gaussian noise. The  
13 SPD differed significantly between all signals ( $p \leq 0.049$  for all comparisons) except between  
14 Lorenz attractor and pink noise. While the brown noise signal had the strongest statistical  
15 persistence, lowest sample entropy, and the highest ENT<sup>1/2</sup> and SPD, the random signal had the  
16 highest SaEn, and the lowest ENT<sup>1/2</sup> and SPD. The Lorenz attractor and pink noise returned  
17 intermediate values for all four analyses.

### 18 **Treadmill and overground walking**

19 The SaEn was significantly greater ( $p=0.033$ ) and the scaling exponent was significantly lower  
20 ( $p=0.012$ ) during treadmill walking compared to overground walking (figure 4A and 4B). There  
21 was no significant difference between the two walking conditions for ENT<sup>1/2</sup> (for overground  
22 walking: mean = 11.3 strides and median = 9 strides and for treadmill walking: mean = 14.6  
23 strides and median = 8 strides) and SPD (for overground walking: mean = 51.6 strides and

1 median = 65 strides and for treadmill walking: mean = 53.0 strides and median = 59.5 strides). It  
2 should be noted that large inter-subject variations existed for  $ENT^{1/2}$  and SPD (figure 4C and  
3 4D). Low non-significant correlations were observed both between SaEn and scaling exponent  
4 (for overground walking:  $R = 0.421$ ,  $R^2 = 0.177$ , Adj.  $R^2 = 0.109$ ,  $p = 0.134$ ; for treadmill  
5 walking:  $R = 0.103$ ,  $R^2 = 0.011$ , Adj.  $R^2 = -0.072$ ,  $p = 0.726$ ) and between  $ENT^{1/2}$  and SPD (for  
6 overground walking:  $R = -0.064$ ,  $R^2 = 0.004$ , Adj.  $R^2 = -0.079$ ,  $p = 0.829$ ; for treadmill walking:  
7  $R = 0.502$ ,  $R^2 = 0.252$ , Adj.  $R^2 = 0.190$ ,  $p = 0.067$ ) (figure 5).

8

9

## 1 Discussion

2 The present study aimed at introducing two novel tools, ENT<sup>1/2</sup> and SPD, as methods to quantify  
3 time dependency in stride-to-stride time intervals during human gait. To validate the use of these  
4 methods, the dynamic characteristics of known theoretical signals (periodic, chaotic, and  
5 random) were assessed by DFA and SaEn in addition to ENT<sup>1/2</sup> and SPD. It was expected that  
6 the periodic signal (brown noise) with low SaEn (high predictability) and with a high scaling  
7 exponent above 0.5 (high statistical persistency) would be characterized with high ENT<sup>1/2</sup> and  
8 SPD, while the random signal (Gaussian noise) with high SaEn (low predictability) and a scaling  
9 exponent close to 0.5 (uncorrelated pattern) would be characterized with low ENT<sup>1/2</sup> and SPD.  
10 Chaotic signals (Lorenz attractor and pink noise) would be characterized by intermediate SaEn,  
11 scaling exponent, ENT<sup>1/2</sup>, and SPD values. This expectation was confirmed which indicated that  
12 the presented methods were able to assess the number of data points involved in creating the  
13 potential time dependency in time series.

14 Furthermore, the present study applied ENT<sup>1/2</sup> and SPD to the stride-to-stride time intervals  
15 recorded during treadmill and overground walking in healthy young adults. Significant lower  
16 predictability and lower statistical persistence were observed during treadmill walking compared  
17 to overground walking. However, no significant difference was observed for ENT<sup>1/2</sup> and SPD  
18 between walking condition.

19 The present study is the first to estimate the time dependency of human gait in an interpretable  
20 scale. The predictability was halved within 11 and 14 consecutive strides during overground and  
21 treadmill walking, respectively, and the statistical persistence was deteriorated within ~50 strides  
22 during both conditions. This indicates a substantial time memory, where information from  
23 previous strides was included in the formation of future strides.

1 During the last twenty years, extensive research has addressed the time dependency and  
2 nonlinear dynamics of the inherent variability in human movements<sup>21,29</sup>. Both basic and applied  
3 research has acknowledged the functional role and importance of the observed movement  
4 variability<sup>22,25,29</sup>. In relation to human gait, variability has been discussed in relation to the  
5 fundamental motor control of walking<sup>7,14,31</sup>, the development of a mature walking pattern<sup>3,15</sup>, and  
6 the impairment following aging<sup>3,13,16</sup>, and pathology<sup>1,9,11,13,18</sup>. The increasing interest in gait  
7 variability has led to a number of different nonlinear tools applied to either kinematic data or  
8 stride characteristics quantifying different characteristics of the time series in question (e.g.  
9 largest Lyapunov exponent quantifying the rate of trajectory divergence or convergence in state  
10 space, approximate and sample entropy quantifying predictability, correlation dimension  
11 quantifying dimensionality, and detrended fluctuation analysis quantifying statistical persistence  
12 or anti-persistence)<sup>28</sup>. While these different tools acknowledge the time dependency in the  
13 investigated time series, they do not quantify this time dependency on an interpretable physical  
14 or physiological time scale.

15 The characteristics and strength of this time dependency in movements observed during walking  
16 could be interpreted as the reliance of the motor control system on previous strides in order to  
17 perform future strides. The present study confirmed previous observations that stride time  
18 intervals during both overground and treadmill walking are characterised by statistical  
19 persistence, meaning that deviations in one direction are statistically likely to be followed by  
20 deviations in the same direction<sup>31</sup>. It has been emphasised that the statistical persistence should  
21 be interpreted within the context of the control process of the parameter in question, the  
22 influence of biomechanical, anatomical, and neuro-muscular redundancy and the task  
23 constraints<sup>6</sup>. Thus, the constraints imposed by the treadmill on the motor control system cause a

1 reduction of the statistical persistence which has been suggested to be linked to a tighter  
2 control<sup>6,31</sup>. Furthermore, Terrier and Deriaz<sup>31</sup> observed that a reduction in statistical persistence  
3 in stride time intervals was accompanied by a reduction in largest Lyapunov exponent of the  
4 centre of mass accelerations during treadmill walking. The authors interpreted this as an increase  
5 in gait stability during treadmill walking compared to overground walking. Interestingly, this  
6 proposed tighter control is accompanied by a lower predictability in the stride time intervals.  
7 Inducing further constraints through use of virtual reality environments with different optic flow,  
8 Katsavelis and colleagues<sup>19</sup> observed an additional decrease in predictability (quantified by  
9 approximate entropy).

10 Two alternative interpretations could be made based on this. It could be speculated that the  
11 tighter control during constraint walking (e.g. treadmill or virtual reality environment) increases  
12 the gait stability (decrease in largest Lyapunov exponent) through more usage of the available  
13 degrees of freedom (increase in sample/approximate entropy). This interpretation suggests that  
14 treadmill walking constitutes an optimal walking condition compared to overground walking.  
15 Alternatively, it could be speculated that the unconstrained overground walking is successfully  
16 performed through a more flexible control which relies on the self-organized interplay of the  
17 degrees of freedom within the body. This interaction creates a movement solution characterized  
18 by higher statistical persistence and a sufficient level of predictability enabling an adaptable  
19 walking pattern. This would furthermore indicate that the human locomotor control during  
20 unconstrained walking is more complex compared to constrained walking as suggested by Costa  
21 et al.<sup>5</sup>. Accordingly, constrained walking could be considered a more challenging task compared  
22 to unconstrained walking for the motor control system to solve which induces a more random-  
23 like pattern in the stride-time intervals. As a consequent, tighter motor control reduces the rate of

1 divergence of the centre of mass accelerations to ensure a stable upper body motion. Although  
2 both explanations are valid, we consider the latter to be the most likely. In support of this  
3 explanation, Dingwell et al.<sup>7</sup> observed that during treadmill walking stride speed fluctuations  
4 exhibited anti-persistence while stride time and stride length exhibited statistical persistence. It  
5 was suggested that while stride speed required a tighter motor control in order to stay in the  
6 middle of the treadmill belt, stride time and stride length was allowed to fluctuate more freely  
7 with more flexible control due to the redundancy of these two parameters<sup>7</sup>. Furthermore, Terrier  
8 et al.<sup>32</sup> observed that during overground walking the stride speed as well as stride time and stride  
9 length exhibited statistical persistence and suggested that these parameters were allowed to  
10 fluctuate freely.

11 In contrast to the scaling exponent and SaEn, the two walking conditions did not induce  
12 differences in ENT<sup>1/2</sup> and SPD. This could indicate that while the constraints imposed by the  
13 treadmill may affect the observed statistical persistence and predictability, it does not seem to  
14 affect the rate at which this deteriorates. Furthermore, the present study investigated the  
15 relationship between SaEn and scaling exponent and between ENT<sup>1/2</sup> and SPD, to assess the  
16 potential shared mechanisms behind the generation of statistical persistence and predictability  
17 observed in stride time intervals. The result showed low non-significant correlations between  
18 SaEn and scaling exponent and between ENT<sup>1/2</sup> and SPD for both walking conditions indicating  
19 no relationship between the generation of statistical persistence predictability and between the  
20 rates of deterioration of the statistical persistence and gradually reduction in predictability. Based  
21 on this, it could be speculated that the sources of these characteristics in the gait patterns does not  
22 changes these parameter synchronously. It is well established that the motor cortex, corticospinal  
23 tract, and spinal cord are involved in the motor control of human locomotion<sup>23</sup>. Although

1 potentially involved in generating the observed time dependency in gait, the actual contribution  
2 from the nervous system is unknown. Furthermore, the need for involvement of higher cortical-  
3 spinal structures in the formation of the statistical persistence observed in human gait has been  
4 questioned<sup>8</sup>. In addition, it has previously been established that the statistical persistence in stride  
5 time intervals is lower in patients of neurological diseases, elderly, and fall prone  
6 individuals<sup>10,13,16</sup>. Thus, pathological and age-related changes in both neurological  
7 musculoskeletal structures could contribute to altered temporal structure of the gait pattern<sup>8</sup>.  
8 However, whether the time dependency is affected in these types of individuals is a topic for  
9 future research.

10 The present study only included walking at the preferred walking speed of the included subjects.  
11 Thus, the motor control was not challenged beyond what could be considered the less demanding  
12 walking task. Thus, it remains unknown if the time dependency of stride time intervals quantified  
13 with  $ENT^{1/2}$  and SPD exhibits the same walking speed relationship as previously shown for  
14 scaling exponent at walking speeds beyond and below the preferred walking speed<sup>17</sup>. Time  
15 dependency was only quantified in the stride-to-stride time intervals. Thus, it remains unknown  
16 if the observed results also apply to other gait characteristics (i.e., stride length, stride speed).  
17 Additionally, future studies should explore the potential of using  $ENT^{1/2}$  and SPD on lower limb  
18 joint angle trajectory data obtained during walking. Previous studies have applied various  
19 nonlinear analyses (e.g. largest Lyapunov exponent, correlation dimension<sup>1,4,26</sup>) to assess joint  
20 angles dynamics. Thus, difference the knee angle joint dynamics has been observed between  
21 young and elderly individuals<sup>4</sup> and between the injured and healthy knee in ACL deficient  
22 patients<sup>30</sup>. It is likely that differences in the time dependency in joint angles during various  
23 locomotion tasks could be detected by  $ENT^{1/2}$  and SPD. The gait pattern of elderly individuals



1 has been observed to be more random compared to younger individuals<sup>4,10</sup>. This would  
2 potentially be characterized by a shorter ENT<sup>1/2</sup> and SPD in the elderly group. However, future  
3 studies should investigate this topic.

4 The present study introduced ENT<sup>1/2</sup> and SPD as novel methods to quantify time dependency  
5 during overground and treadmill walking and were able to show that predictability in stride time  
6 intervals was halved after approximate 14 strides and that the statistical persistence was  
7 deteriorated to uncorrelated noise after approximately 50 strides. These observations were  
8 accompanied by a lower statistical persistence and lower predictability during treadmill walking  
9 compared to overground walking.

## 10 **Acknowledgements**

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12 Health (P20 GM109090).

## References

1. Alkjaer T., P. C. Raffalt, H. Dalsgaard, E. B. Simonsen, N. C. Petersen, H. Bliddal and M. Henriksen. Gait variability and motor control in people with knee osteoarthritis. *Gait Posture* 42: 479-484, 2015.
2. Baltich J., V. Von Tscharner, P. Zandiyeh and B. M. Nigg. Quantification and reliability of center of pressure movement during balance tasks of varying difficulty. *Gait Posture* 40: 327-332, 2014.
3. Bisi M. C. and R. Stagni. Complexity of human gait pattern at different ages assessed using multiscale entropy: From development to decline. *Gait Posture* 47: 37-42, 2016.
4. Buzzi U. H., N. Stergiou, M. J. Kurz, P. A. Hageman and J. Heidel. Nonlinear dynamics indicates aging affects variability during gait. *Clin Biomech (Bristol, Avon)* 18: 435-443, 2003.
5. Costa M., C. K. Peng, A. L. Goldberger and J. M. Hausdorff. Multiscale entropy analysis of human gait dynamics. *Physica A* 330: 53-60, 2003.
6. Dingwell J. B. and J. P. Cusumano. Re-interpreting detrended fluctuation analyses of stride-to-stride variability in human walking. *Gait Posture* 32: 348-353, 2010.
7. Dingwell J. B., J. John and J. P. Cusumano. Do humans optimally exploit redundancy to control step variability in walking? *PLoS Comput Biol* 6: e1000856, 2010.
8. Gates D. H., J. L. Su and J. B. Dingwell. Possible Biomechanical Origins of the Long-Range Correlations in Stride Intervals of Walking. *Physica A* 380: 259-270, 2007.
9. Georgoulis A. D., C. Moraiti, S. Ristanis and N. Stergiou. A novel approach to measure variability in the anterior cruciate ligament deficient knee during walking: the use of the approximate entropy in orthopaedics. *J Clin Monit Comput* 20: 11-18, 2006.
10. Goldberger A. L., L. A. Amaral, J. M. Hausdorff, P. Ivanov, C. K. Peng and H. E. Stanley. Fractal dynamics in physiology: alterations with disease and aging. *Proc Natl Acad Sci U S A* 99 Suppl 1: 2466-2472, 2002.
11. Hausdorff J. M. Gait dynamics in Parkinson's disease: common and distinct behavior among stride length, gait variability, and fractal-like scaling. *Chaos* 19: 026113, 2009.
12. Hausdorff J. M., Y. Ashkenazy, C. K. Peng, P. C. Ivanov, H. E. Stanley and A. L. Goldberger. When human walking becomes random walking: fractal analysis and modeling of gait rhythm fluctuations. *Physica A* 302: 138-147, 2001.
13. Hausdorff J. M., S. L. Mitchell, R. Firtion, C. K. Peng, M. E. Cudkowicz, J. Y. Wei and A. L. Goldberger. Altered fractal dynamics of gait: reduced stride-interval correlations with aging and Huntington's disease. *J Appl Physiol (1985)* 82: 262-269, 1997.
14. Hausdorff J. M., P. L. Purdon, C. K. Peng, Z. Ladin, J. Y. Wei and A. L. Goldberger. Fractal dynamics of human gait: stability of long-range correlations in stride interval fluctuations. *J Appl Physiol (1985)* 80: 1448-1457, 1996.
15. Hausdorff J. M., L. Zeman, C. Peng and A. L. Goldberger. Maturation of gait dynamics: stride-to-stride variability and its temporal organization in children. *J Appl Physiol (1985)* 86: 1040-1047, 1999.
16. Herman T., N. Giladi, T. Gurevich and J. M. Hausdorff. Gait instability and fractal dynamics of older adults with a "cautious" gait: why do certain older adults walk fearfully? *Gait Posture* 21: 178-185, 2005.
17. Jordan K., J. H. Challis and K. M. Newell. Walking speed influences on gait cycle variability. *Gait Posture* 26: 128-134, 2007.
18. Kaipust J. P., J. M. Huisinga, M. Filipi and N. Stergiou. Gait variability measures reveal differences between multiple sclerosis patients and healthy controls. *Motor Control* 16: 229-244, 2012.
19. Katsavelis D., M. Mukherjee, L. Decker and N. Stergiou. The effect of virtual reality on gait variability. *Nonlinear Dynamics Psychol Life Sci* 14: 239-256, 2010.
20. Lee S. J. and J. Hidler. Biomechanics of overground vs. treadmill walking in healthy individuals. *J Appl Physiol (1985)* 104: 747-755, 2008.

- 1 21. Newell K. M. and D. M. Corcos. Issues in variability and motor control. In: *Variability and motor*  
2 *control*, edited by K. M. Newell and D. M. Corcos. Champagne, IL: Human Kinetics, 1993.
- 3 22. Newell K. M. and D. E. Vaillancourt. Dimensional change in motor learning. *Hum Mov Sci* 20:  
4 695-715, 2001.
- 5 23. Nielsen J. B. How we walk: central control of muscle activity during human walking.  
6 *Neuroscientist* 9: 195-204, 2003.
- 7 24. Peng C. K., S. Havlin, H. E. Stanley and A. L. Goldberger. Quantification of scaling exponents and  
8 crossover phenomena in nonstationary heartbeat time series. *Chaos* 5: 82-87, 1995.
- 9 25. Preatoni E., J. Hamill, A. J. Harrison, K. Hayes, R. E. Van Emmerik, C. Wilson and R. Rodano.  
10 Movement variability and skills monitoring in sports. *Sports Biomech* 12: 69-92, 2013.
- 11 26. Raffalt P. C., M. K. Guul, A. N. Nielsen, S. Puthusserypady and T. Alkjaer. Economy, Movement  
12 Dynamics, and Muscle Activity of Human Walking at Different Speeds. *Sci Rep* 7: 43986, 2017.
- 13 27. Richman J. S. and J. R. Moorman. Physiological time-series analysis using approximate entropy  
14 and sample entropy. *Am J Physiol Heart Circ Physiol* 278: H2039-2049, 2000.
- 15 28. Stergiou N. *Nonlinear analysis for human movement variability*. Boca Raton, FL: CRC Press, 2016.
- 16 29. Stergiou N. and L. M. Decker. Human movement variability, nonlinear dynamics, and pathology:  
17 is there a connection? *Hum Mov Sci* 30: 869-888, 2011.
- 18 30. Stergiou N., C. Moraiti, G. Giakas, S. Ristanis and A. D. Georgoulis. The effect of the walking  
19 speed on the stability of the anterior cruciate ligament deficient knee. *Clin Biomech (Bristol, Avon)* 19:  
20 957-963, 2004.
- 21 31. Terrier P. and O. Deriaz. Kinematic variability, fractal dynamics and local dynamic stability of  
22 treadmill walking. *J Neuroeng Rehabil* 8: 12, 2011.
- 23 32. Terrier P., V. Turner and Y. Schutz. GPS analysis of human locomotion: further evidence for long-  
24 range correlations in stride-to-stride fluctuations of gait parameters. *Hum Mov Sci* 24: 97-115, 2005.
- 25 33. Yentes J. M., N. Hunt, K. K. Schmid, J. P. Kaipust, D. McGrath and N. Stergiou. The appropriate  
26 use of approximate entropy and sample entropy with short data sets. *Ann Biomed Eng* 41: 349-365,  
27 2013.
- 28 34. Zandiyeh P. and V. Von Tscherner. Reshape scale method: a novel multi scale entropic analysis  
29 approach. *Physica A* 392: 6265-6272, 2013.

30

31

1 **Legends**

2 **Table 1:** Mean  $\pm$  SD of scaling exponent, sample entropy, entropic half-life and statistical persistence  
3 decay of twenty iteration of brown noise, Lorenz attractor, pink noise signal and white Gaussian noise.  
4 NOTE: a) indicates significant difference from brown noise signal, b) indicates significance different  
5 from Lorenz attractor signal, c) indicates significant difference from pink noise signal ( $p < 0.05$ ).

6

7 **Figure 1:** Examples of the brown noise signal, Lorenz attractor signal, pink noise signal, and the white  
8 Gaussian noise signal (top four signals) and an example of the stride time interval time series for  
9 overground and treadmill walking (bottom two signals). One hundred data points or stride time intervals  
10 are depicted.

11

12 **Figure 2:** Calculation procedure for  $ENT^{1/2}$  (see text for details).

13

14 **Figure 3:** Calculation procedure for SPD (see text for details).

15

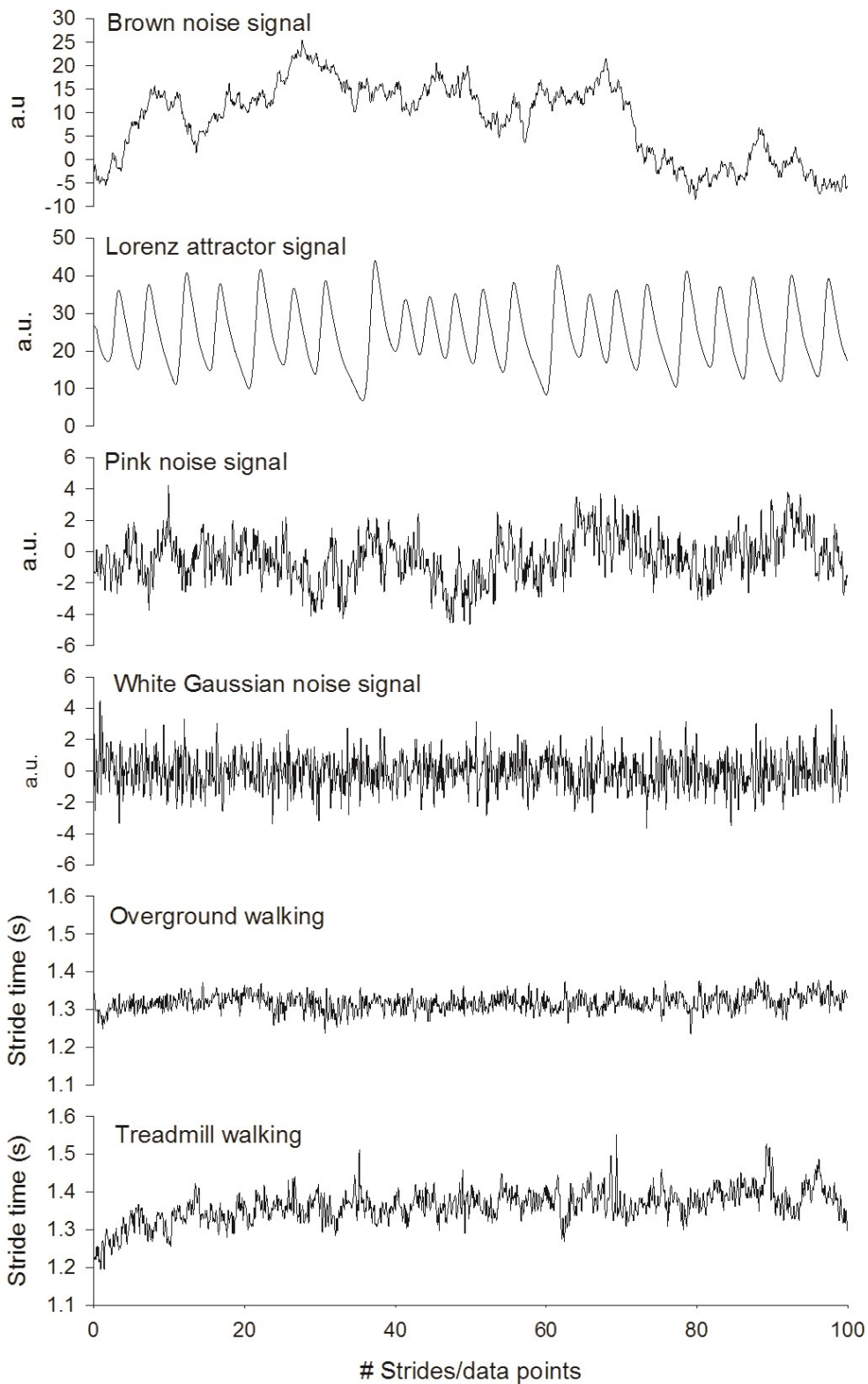
16 **Figure 4A-D:** Boxplot including group mean (dashed line) and median (solid line) for SaEn (A), scaling  
17 exponent (B),  $ENT^{1/2}$  (C), and SPD (D) for overground walking (OG) and treadmill walking (TM).

18

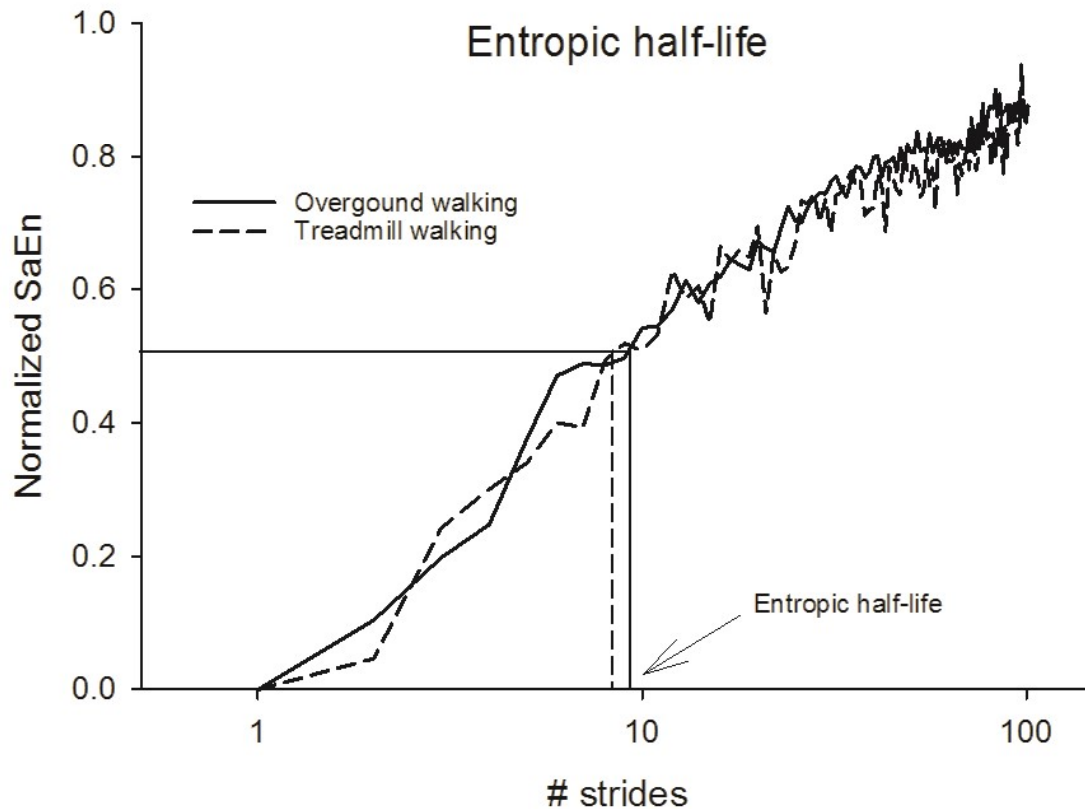
19 **Figure 5:** Linear regression of SaEn and scaling exponent and of  $ENT^{1/2}$  and SPD for both overground  
20 and treadmill walking.

**Table 1:** Mean  $\pm$  SD of scaling exponent, sample entropy, entropic half-life and statistical persistence decay of twenty iteration of brown noise, Lorenz attractor, pink noise signal and white Gaussian noise. NOTE: a) indicates significant difference from brown noise signal, b) indicates significance different from Lorenz attractor signal, c) indicates significant difference from pink noise signal ( $p < 0.05$ ).

	Brown noise	Lorenz attractor	Pink noise	Gaussian noise
Alpha	$1.40 \pm 0.05$	$1.08 \pm 0.03^a$	$0.93 \pm 0.06^{a,b}$	$0.47 \pm 0.03^{a,b,c}$
SaEn	$0.18 \pm 0.08$	$0.22 \pm 0.01$	$1.80 \pm 0.05^{a,b}$	$2.18 \pm 0.01^{a,b,c}$
ENT $\frac{1}{2}$	$43.30 \pm 29.88$	$8.00 \pm 0.00^a$	$4.20 \pm 0.83^{a,b}$	$2.60 \pm 0.82^{a,b}$
SPD	$68.65 \pm 11.14$	$16.95 \pm 15.10^a$	$45.05 \pm 18.59^a$	$1.05 \pm 0.22^{a,b,c}$

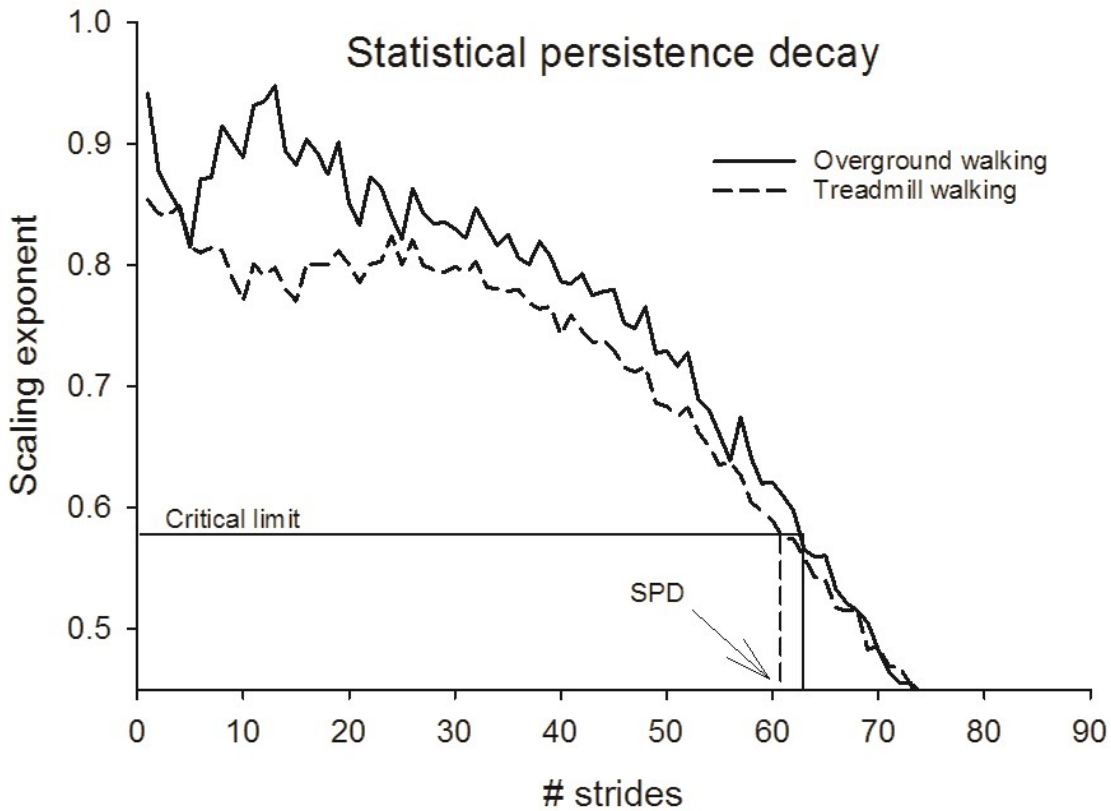


- 1) Calculate SaEn for the original time series ( $SaEn_{OR}$ )
- 2) Create one hundred reshaped time series through successively reshaping the original time series which gradually randomised each new reshaped time series following the principle described below:  
 Original time series: [1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16]  
 Reshaped time series 2: [1 3 5 7 9 11 13 15 2 4 6 8 10 12 14 16]  
 Reshaped time series 3: [1 4 7 10 13 16 2 5 8 11 14 3 6 9 12 15]  
 Reshaped time series 4: [1 5 9 13 2 6 10 14 3 7 11 15 4 8 12 16]
- 3) Calculate SaEn for each reshape time series ( $SaEn_{RS}$ )
- 4) Calculate the average SaEn for one hundred randomized time series created by a random permutation of the data points in the original time series ( $SaEn_{RAN}$ )
- 5) Normalize the  $SaEn_{RS}$  to the difference between the  $SaEn_{OR}$  and  $SaEn_{RAN}$  according to equation 2 (see text)
- 6) Plot the  $SaEn_{RS}$  in as a function of the stride number (see below)
- 7) The stride number at which  $SaEn_{RS}$  exceeds 0.5 is defined as the entropic half-life

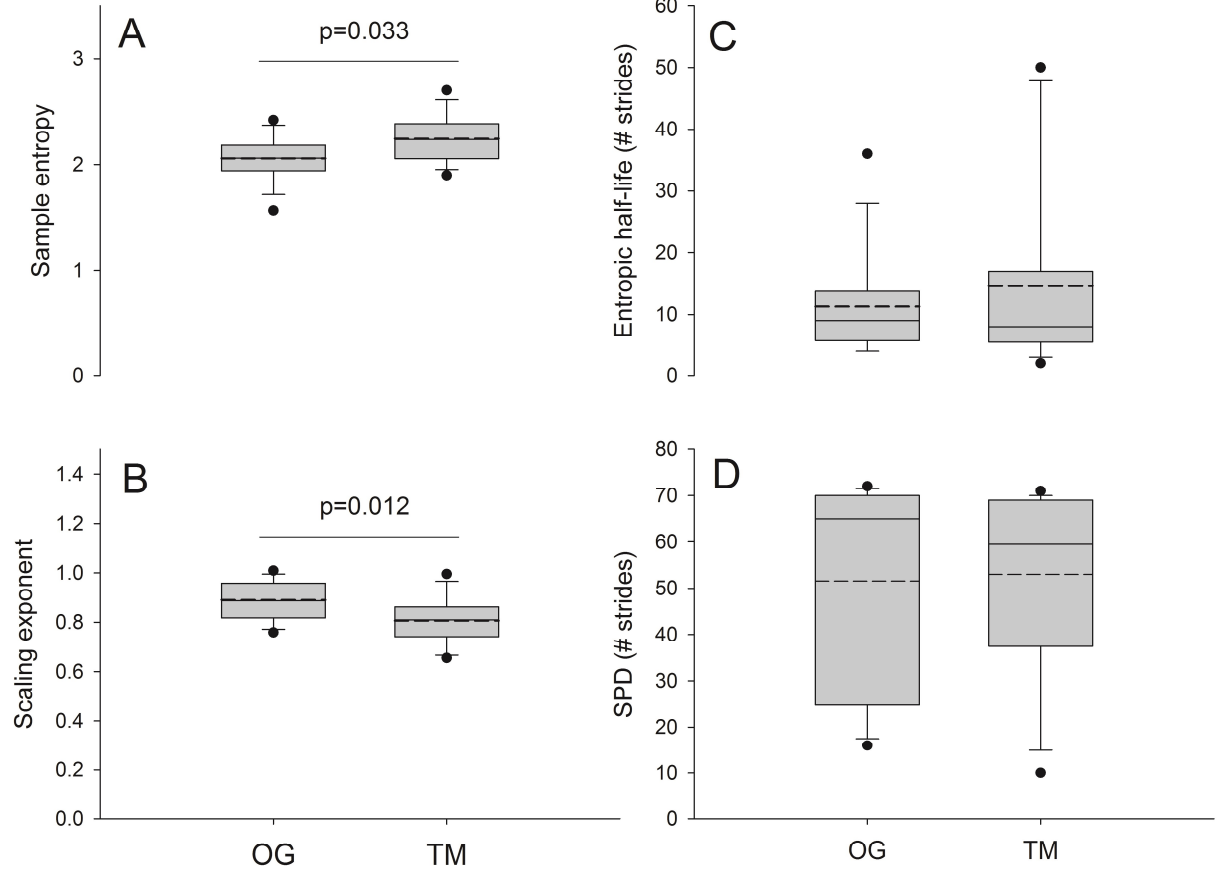


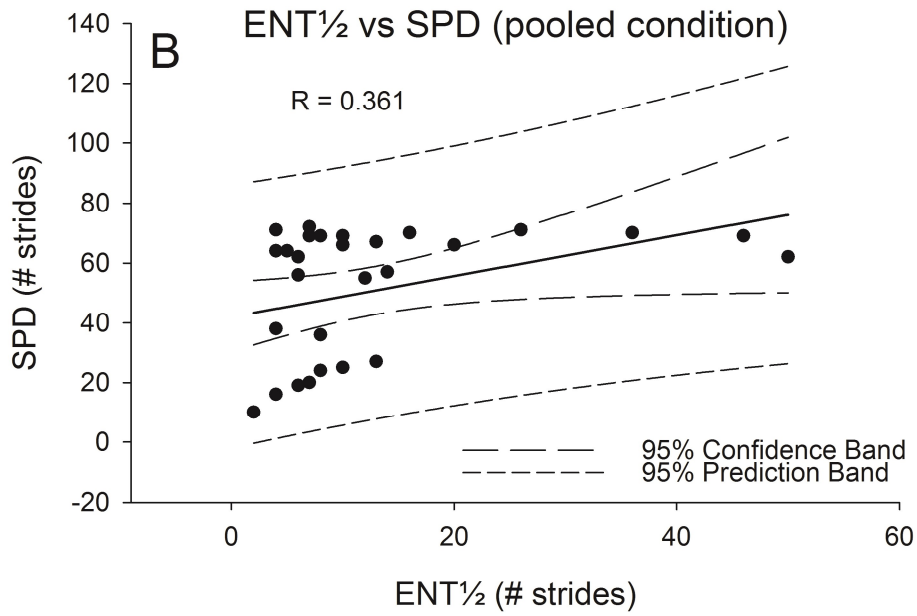
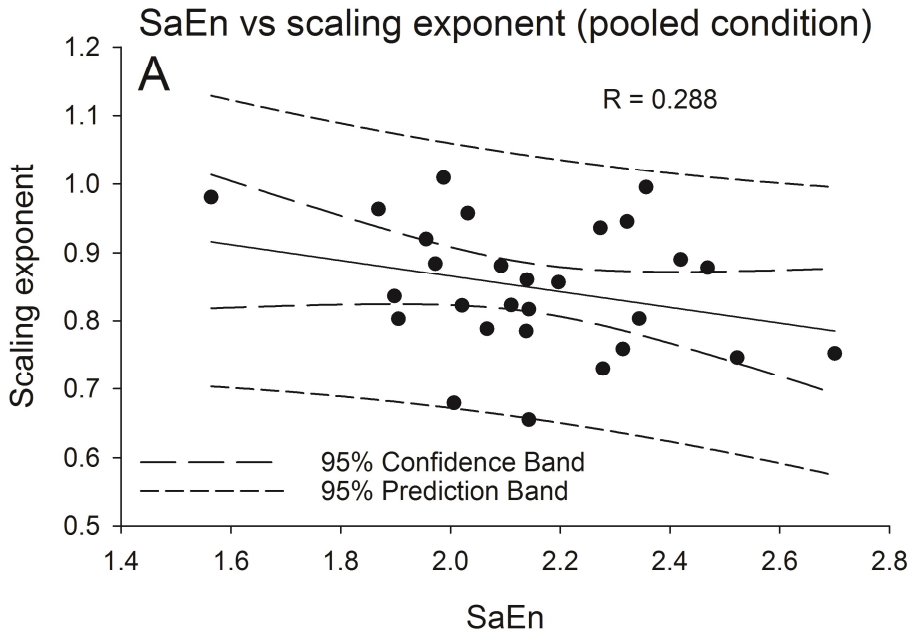
- 1) Calculate scaling exponent for the original time series
- 2) Create one hundred reshaped time series through successively reshaping the original time series which gradually randomised each new reshaped time series following the principle described below:
 

Original time series:	[1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16]
Reshaped time series 2:	[1 3 5 7 9 11 13 15 2 4 6 8 10 12 14 16]
Reshaped time series 3:	[1 4 7 10 13 16 2 5 8 11 14 3 6 9 12 15]
Reshaped time series 4:	[1 5 9 13 2 6 10 14 3 7 11 15 4 8 12 16]
- 3) Calculate scaling exponent for each reshaped time series
- 4) Calculate the average and standard deviation of the scaling exponents of one hundred randomized time series created by a random permutation of the data points in the original time series ( $\mu_{\text{aRAN}}$  and  $\sigma_{\text{aRAN}}$ )
- 5) Calculate a critical limit according to equation 5 (see text)
- 6) Plot the scaling exponents of the reshaped time series as function of the stride number (see below)
- 7) The stride number at which the scaling exponent is below the critical limit is defined as the statistical persistence decay (SPD)









## **SUPPLEMENTARY MATERIAL**

### **Title:**

Introducing Statistical Persistence Decay – A Quantification of Stride-to-Stride Time Interval  
Dependency in Human Gait

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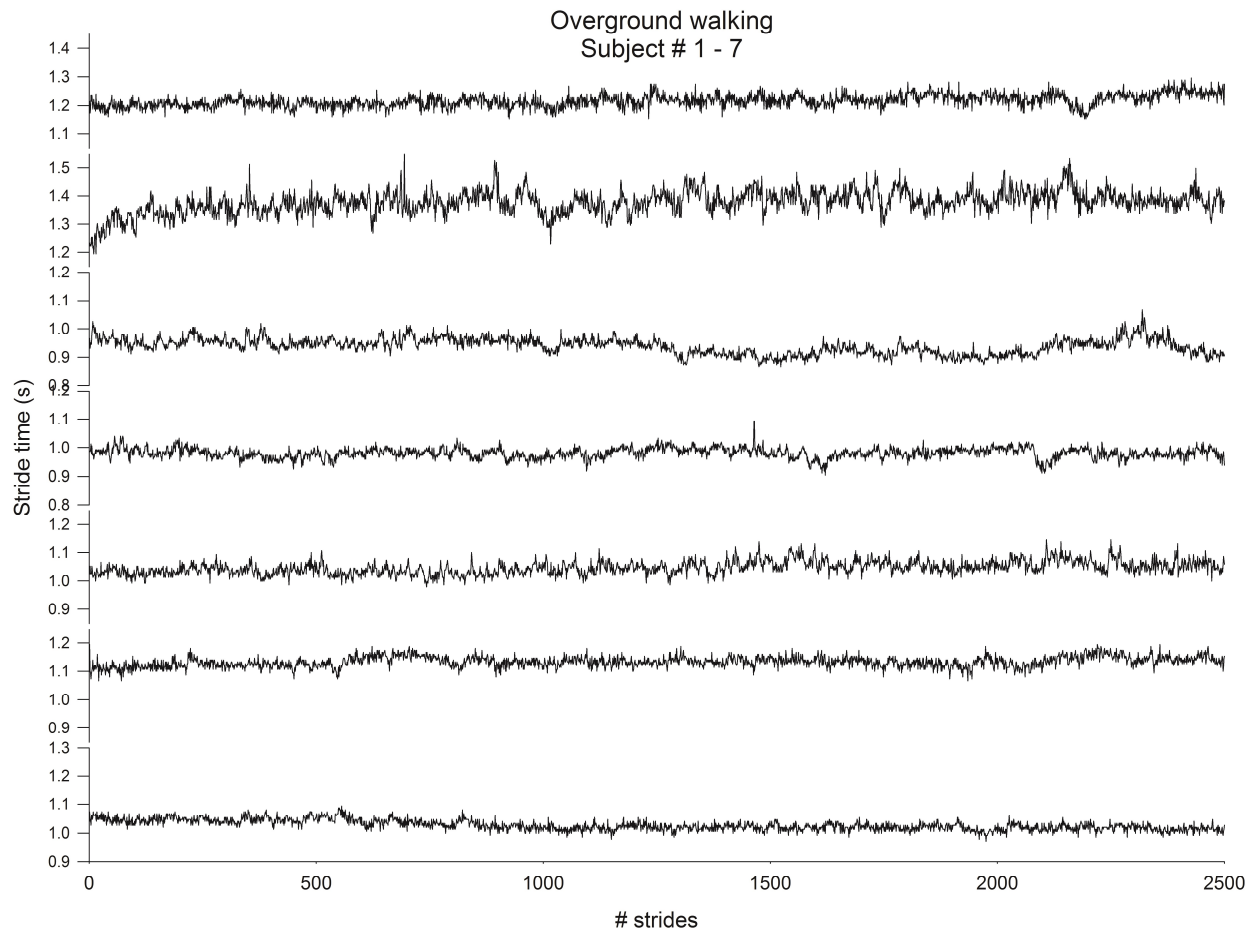
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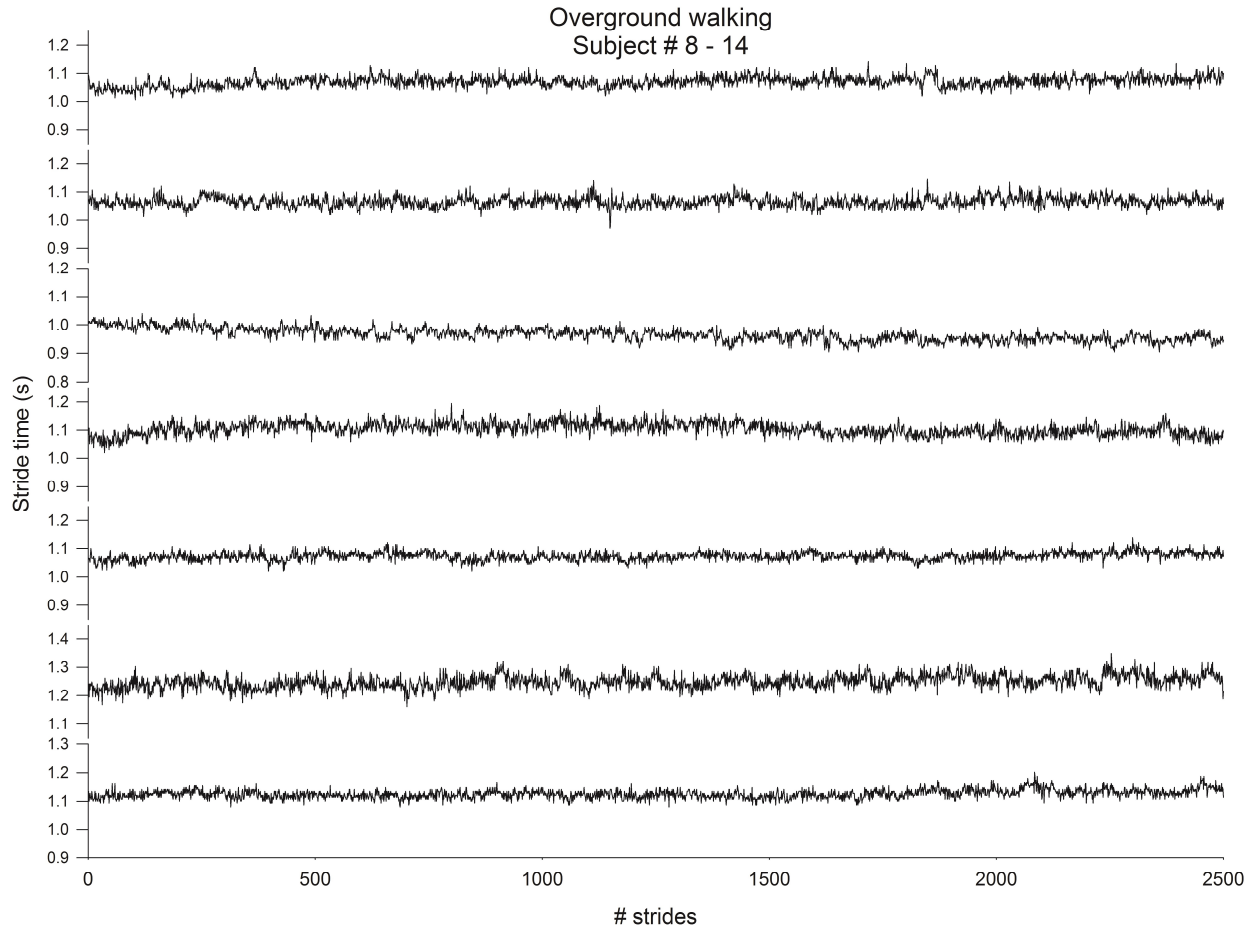
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## Supplementary material 1

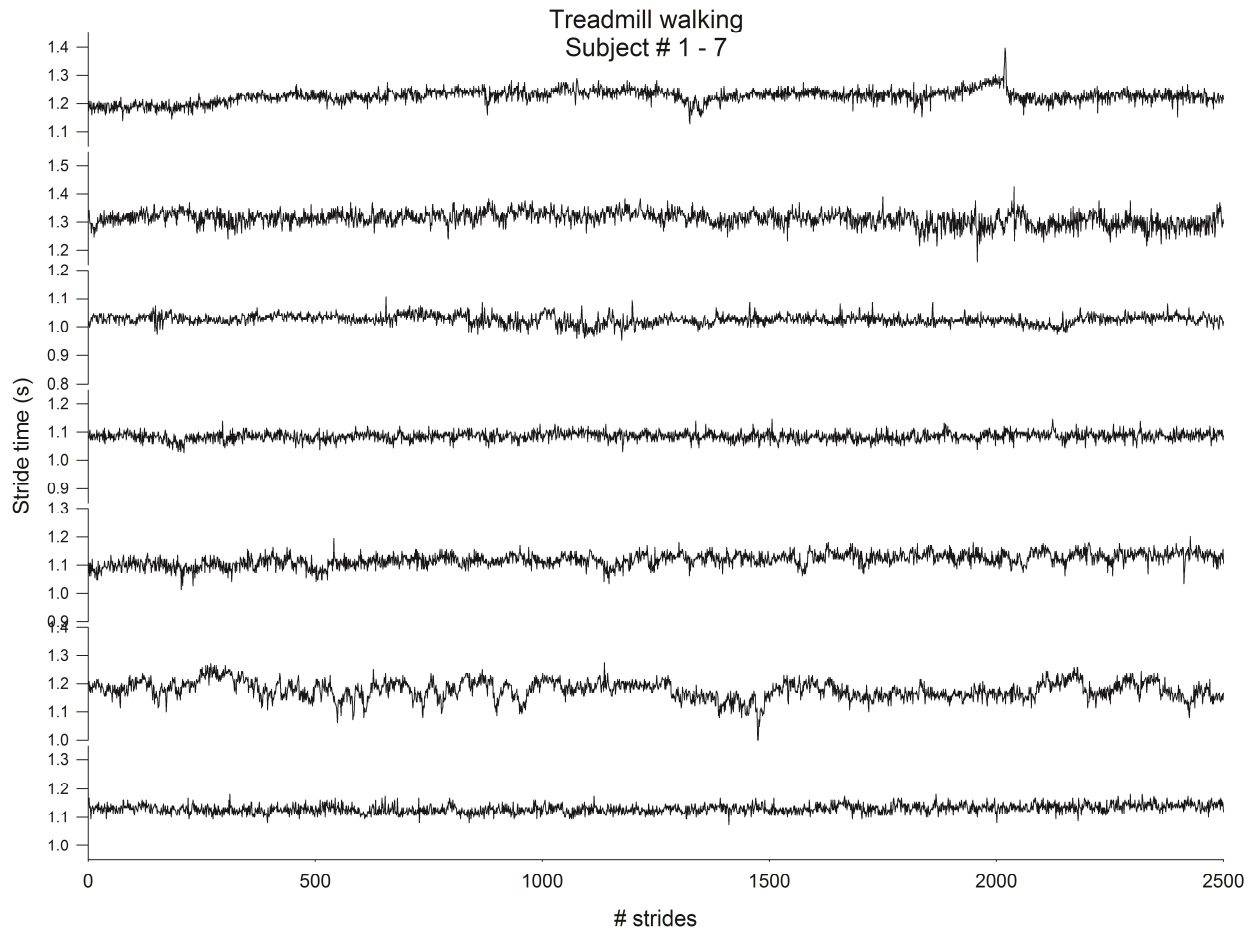
In this material the stride time interval time series for each subject during overground and treadmill walking is presented. The trials included 2500 strides and are presented in four figures.



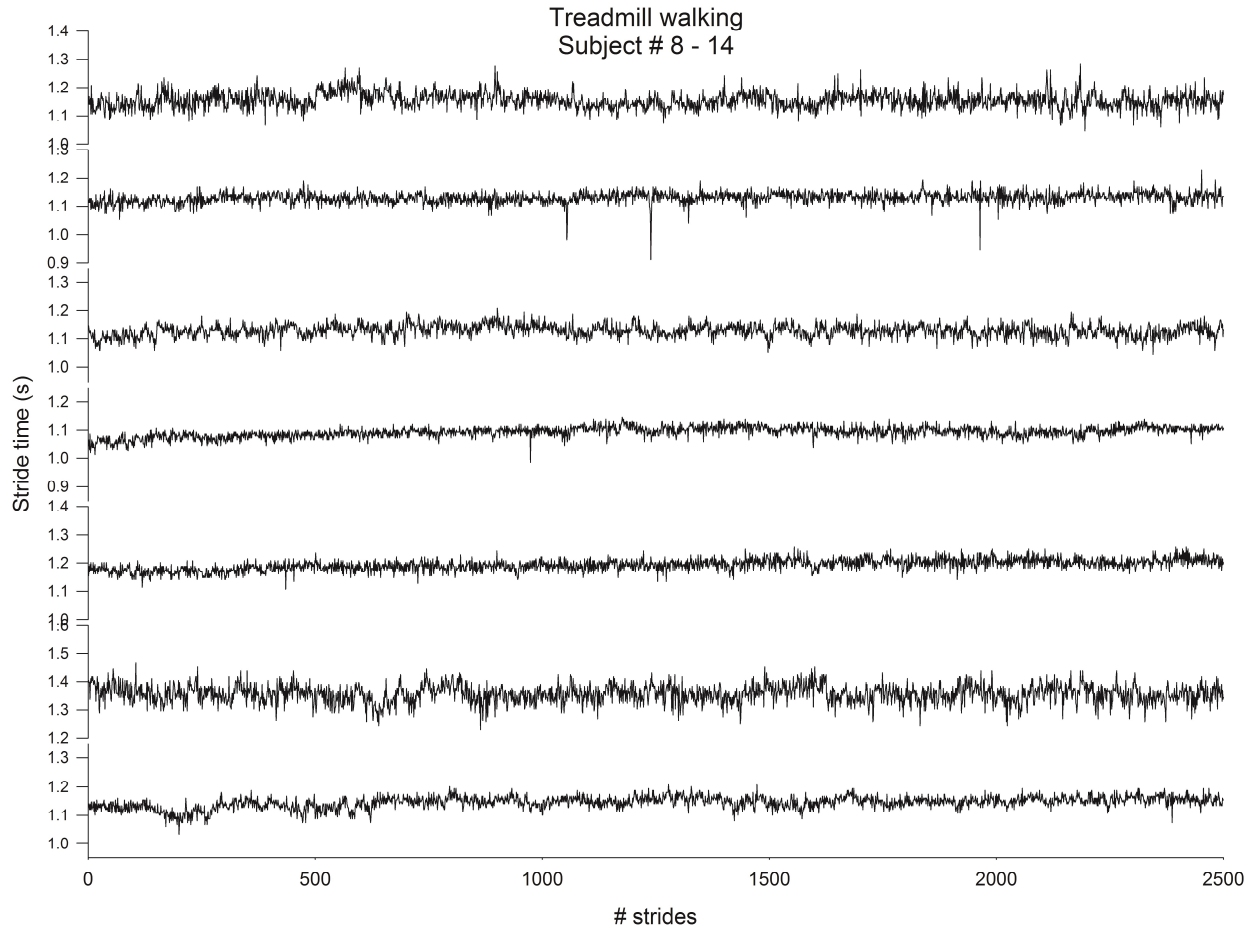
**Figure S1.1:** Stride time interval time series for subjects 1 – 7 during overground walking.



**Figure S1.2:** Stride time interval time series for subjects 8 – 14 during overground walking.



**Figure S1.3:** Stride time interval time series for subjects 1 – 7 during treadmill walking.



**Figure S1.4:** Stride time interval time series for subjects 8 – 14 during treadmill walking.

## Supplementary material 2

### *Introduction*

Sample entropy (SaEn) analysis of a time series require the determination of three parameters 1)  $r$ , the similarity criterion, 2)  $m$ , the length of data that will be compared and 3)  $N$ , the length of the time series prior to the calculation. As shown by Yentes and colleagues (2013), the selected parameters are crucial to the outcome of sample entropy (SaEn) calculations. Applying different parameters to the same time series will potentially change the outcome. To avoid this, the use of SaEn to time series should be accompanied by a test for parameter consistency. The purpose of the supplementary material was to investigate the parameter consistency for the time series in the present study.

### *Method*

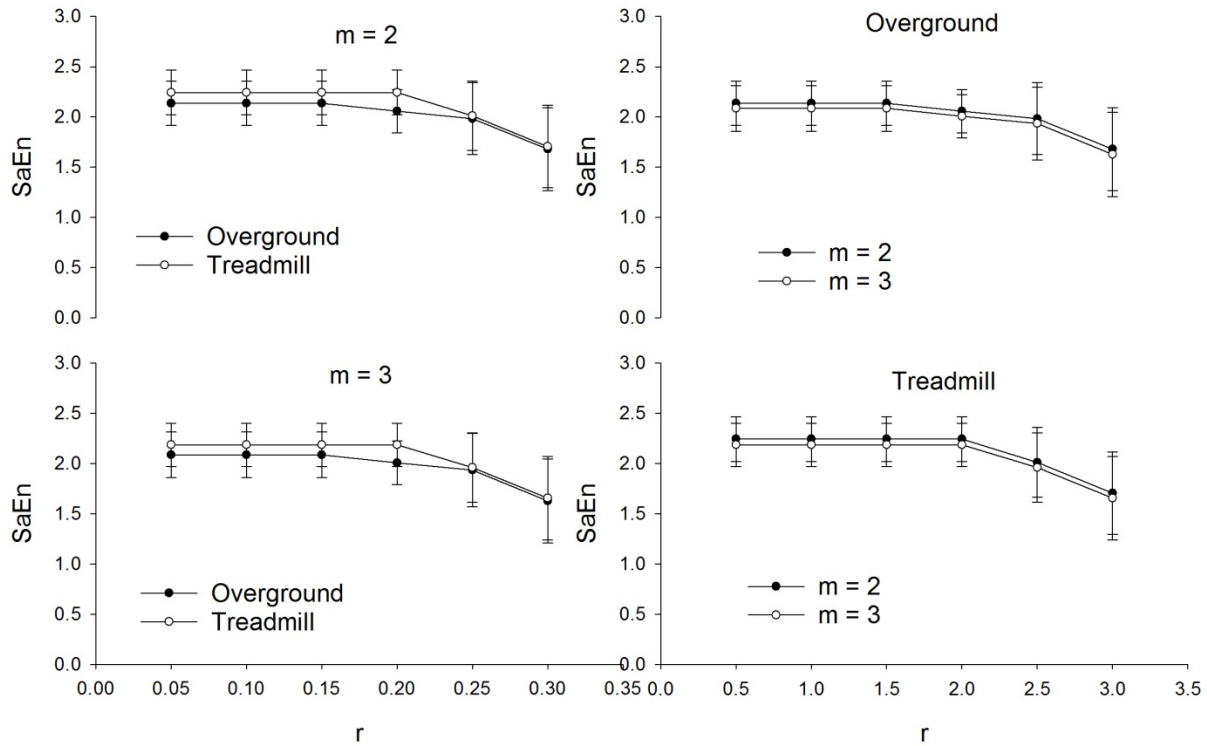
The present study used 2500 data point in each time series which is well beyond what is needed to achieve a consistent SaEn outcome (Yentes et al. 2013). Therefore, parameter consistency analysis was only performed for parameters  $r$  and  $m$ . SaEn was calculated for the stride time interval time series during both treadmill and overground walking using  $r = 0.5, 1, 1.5, 2, 2.5$  and  $3$  and  $m = 2$  and  $3$ . A two-way ANOVA for repeated measures was applied to evaluate the effect of different  $r$  and  $m$  values for both the overground and treadmill data. In case of significant effects, a Holm-Sidak post hoc test was completed. Level of significance was set at 5 %. Calculations were conducted in SigmaPlot (Systat Software, Inc. 2014, version 13.0, Germany).

### *Results and Conclusion*

There was no significant effect of the different  $r$  values on the SaEn values (figure S1). The non-significant decrease in SaEn at higher  $r$  values was similar for both the overground and treadmill walking data. There was a significant main effect of walking condition ( $p=0.029$  for  $m=2$  and  $p=0.036$  for  $m=3$ ) indicating that across  $r$  values the SaEn was significantly higher during treadmill walking compared to overground walking. There was a significant effect of the parameter  $m$  for both overground walking ( $p<0.001$ ) and treadmill walking ( $p<0.001$ ) (figure S1). Thus, SaEn was significantly higher with  $m=2$  compared to  $m=3$  for both walking conditions. It is noteworthy that the inter-condition relationship did not change with changes in  $r$ . Equally, the inter-parameter relationship of  $m$  did not change with changes in  $r$ .



Based on these results (figure S2.1), it could be concluded that there in general was a satisfying parameter consistency for the tested parameters and that  $r = 0.2$  and  $m = 2$  was well suited for SaEn calculations of the data in the present study.



**Figure S2.1:** Group mean  $\pm$  SD of SaEn calculated for overground and treadmill walking with  $r$  values of 0.05, 0.1, 0.15, 0.2, 0.25 and 0.3 and  $m$  values of 2 and 3.

### References

Yentes, J. M., Hunt, N., Schmid, K. K., Kaipust, J. P., McGrath, D., & Stergiou, N. (2013). The appropriate use of approximate entropy and sample entropy with short data sets. *Ann Biomed Eng*, 41, 349-365.