

## Original Article

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**Author for correspondence:**

L. C. M. Vanderlei, Department of Physiotherapy, Sao Paulo State University, UNESP, Rua Roberto Simonsen, 305 – Centro Educacional, Presidente Prudente 19060-900, Brazil.  
Tel: +55 (18) 3229-5388; Fax: +55 (18) 3229-5389; E-mail: [lcm.vanderlei@unesp.br](mailto:lcm.vanderlei@unesp.br)

# Non-linear regulation of cardiac autonomic modulation in obese youths: interpolation of ultra-short time series

David M. Garner<sup>1</sup>, Franciele M. Vanderlei<sup>2</sup> , Vitor E. Valenti<sup>3</sup>  and Luiz Carlos M. Vanderlei<sup>2</sup>

<sup>1</sup>Cardiorespiratory Research Group, Department of Biological and Medical Sciences, Faculty of Health and Life Sciences, Oxford Brookes University, Oxford, UK; <sup>2</sup>Department of Physiotherapy, Sao Paulo State University, UNESP, Presidente Prudente, Brazil and <sup>3</sup>Autonomic Nervous System Center, Sao Paulo State University, UNESP, Presidente Prudente, Brazil

**Abstract**

**Background:** In this study, we applied ultra-short time series of interbeat intervals (RR-intervals) to evaluate heart rate variability through default chaotic global techniques with the purpose of discriminating obese youths from non-obese youth patients. **Method:** Chaotic global analysis of the RR-intervals from the electrocardiogram and pre-processing adjustments was undertaken. The effect of cubic spline interpolations was assessed, while the spectral parameters remained fixed. Exactly, 125 RR-intervals of data were recorded. **Results:** CFP1, CFP3, and CFP6 were the only significant combinations of chaotic globals when the standard conditions were enforced and at the level  $p < 0.01$  (or  $< 1\%$ ). These significances were achieved via Kruskal–Wallis and Cohen’s  $d_s$  effects sizes tests of significance after Anderson–Darling and Lilliefors statistical tests indicated non-normal distributions in the majority of cases. Adjustments of the cubic spline interpolation from 1 to 13 Hz were revealed to be inconsequential when measured by Kruskal–Wallis and Cohen’s  $d_s$ , regarding the outcome between the two datasets. **Conclusion:** Chaotic global analysis was offered as a robust technique to distinguish autonomic dysfunction in obese youths. It can discriminate the two different groups using ultra-short data lengths, and no cubic spline interpolations need be applied.

Heart rate variability is an important sign for diagnosing cardiac pathological states.<sup>1</sup> Mathematical algorithms founded on non-linear dynamics are useful when analysing these conditions. They are vital when developing new methods to reach an early differential diagnosis about cardiovascular disease conditions.<sup>2</sup> The sympathetic and parasympathetic nervous systems’ connections have been demonstrated to influence heart rate variability by non-linear neurological cross-talk. Heart rate variability is a simple, inexpensive, and non-invasive way of monitoring the cardiac branches of the Autonomic Nervous System. Other procedures can be unresponsive such as with Sympathetic Skin Response<sup>3</sup> or, too complicated and highly priced as with Quantitative Pupillography.<sup>4</sup>

The beat of electrocardiographic interbeat intervals (RR-intervals) derived from the PQRST-motif can pulsate in an irregular and often chaotic manner.<sup>5,6</sup> This has been previously undertaken,<sup>7,8</sup> but with much longer data sets (1000 RR-intervals) and without judging the potential consequences of cubic spline interpolations.<sup>9</sup> These chaotic global metrics are especially sensitive to unpredictabilities. This is predominant when compared with those based on linear descriptive statistics, conventional non-linear, or geometric routines. The larger the response, usually the healthier the patients’ physiological status. Less chaos can typically be interpreted as a mathematical marker for dynamical disease states, in particular.<sup>10</sup> *Dynamical diseases* are characterised by unexpected instabilities in the qualitative dynamics of physiological processes. This leads to irregular dynamics and then *pathological states*. So, there is a physiological connection between non-linear dynamics (or complexity theory) and clinical medicine.<sup>11</sup> This method is valuable to the clinical team to identify subtle changes in the Autonomic Nervous System, and to predict the risk of problems.

Such computations are advantageous when assessing surgical patients<sup>12</sup> principally anaesthetised<sup>13</sup> or unable to communicate distress such as in sleep apnea patients<sup>14</sup> or those experiencing “air hunger.”<sup>15,16</sup> However, the use of ultra-short time series of RR-intervals to evaluate heart rate variability through default chaotic global techniques for evaluation of health conditions is unknown in the literature.

Through the RR-intervals, we compute three chaotic global parameters and seven groupings to determine the control from the experimental time series. We assumed that these patients

from obese youths' datasets presented autonomic alterations that can be observed with the proposed analyses.

## Methods

Patient selection and assessments were exactly as with the studies by Vanderlei et al<sup>7</sup> and Garner et al.<sup>8</sup> All procedures performed in studies involving human patients were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. All volunteers signed a consent letter and were informed of the procedures and objectives of the study. The study's procedures were all approved by the Research Ethics Committee of Sao Paulo State University, UNESP (Number Protocol No. 11/2011).

### Chaotic globals and chaotic forward parameters

There are three types of chaotic global parameter. They are characterised as *high spectral* Entropy,<sup>7</sup> *high spectral* Detrended Fluctuation Analysis,<sup>7</sup> and Spectral Multi-Taper Method.<sup>17</sup> All are functions of the Multi-Taper Method power spectrum.<sup>18</sup> As such, Multi-Taper Method is a form of power spectrum that has been revealed to be beneficial for spectral estimation.<sup>19</sup> Its major advantage is the minimization of spectral leakage. Functions described as discrete prolate spheroidal sequences, often referred to as Slepian Sequences,<sup>20</sup> are a set of functions that optimise their windows. Regarding *high spectral* Entropy, Shannon entropy<sup>21</sup> is applied directly onto the Multi-Taper Method power spectrum. Whereas with *high spectral* Detrended Fluctuation Analysis, Peng et al's algorithm, the Detrended Fluctuation Analysis<sup>22</sup> is applied directly onto an identical Multi-Taper Method power spectrum. Spectral Multi-Taper Method<sup>17</sup> is dependent on elevated broadband noise intensities generated in Multi-Taper Method power spectra by irregular and often chaotic signals. When broadband noise is increased significantly during an elevated chaotic response, the area beneath the power spectrum increases. Spectral Multi-Taper Method is the area between this power spectrum and the baseline.

Chaotic Forward Parameters 1–7 (CFP1–CFP7)<sup>19</sup> are enforced on the electrocardiographics RR-intervals for the non-obese and obese youths' patients. *High spectral* Detrended Fluctuation Analysis responds to levels of chaos contrarily to the others, so we deduct its value from unity. Weightings of unity are stated here for each of the three chaotic global parameters.

There are seven non-trivial combinations of three chaotic global values.<sup>23</sup> It is anticipated that CFP1 that applies all three should be the most statistically robust. This for the reason it takes the information and processes it in three different ways. The summation of the three would be expected to deviate greater than single or double permutations. The potential danger is since we are only computing spectral components, phase information is lost.

### Principal component analysis

Principal component analysis<sup>24,25</sup> is a multivariate statistical technique for analysing the complexity of high-dimensional data sets. Principal component analysis is useful when sources of variability in the data need to be explained or reducing the complexity of the data and through this assess the data with lesser dimensions. The primary aim of principal component analysis is to represent the data with fewer variables, while sustaining the majority of the total variance. There are two major properties of the

principal component analysis. First, the technique is non-parametric so no prior knowledge can be incorporated. And then secondly, principal component analysis data reduction often incurs a loss of information.

Next, there are the assumptions of the technique. Initially linearity, this accepts the data set to be linear combinations of the variables. Then, the importance of mean and covariance, hence no assurance that the direction of maximum variance, will contain good discriminative features. And finally, large variance has the most important dynamics and the lowest corresponding to noise.

When interpreting the principal component analysis, four points should be considered. First, the higher the component loadings the more important that the variable is to the component. Second, positive and negative loadings are interpreted as mixed. Third, the specific sign of these mixed loadings is unimportant. Finally, the rotated component matrix is vital.

### Effect sizes by Cohen's $d_s$

Cohen's  $d$ <sup>26,27</sup> generally denotes the entire group termed effect sizes. To quantify the magnitude of difference between protocols for significant differences, the effect size was estimated through a sub-group Cohen's  $d_s$ .<sup>28</sup> Cohen's  $d_s$  represents the standardised mean difference of an effect. It can be used to calculate effects across studies even when the dependent variables are measured in alternative ways or even when completely different measures are used. It varies from zero to infinity, which may be positive or negative. However, Cohen refers to the standardised mean difference between two groups of independent observations for the appropriate *sample* as  $d_s$ .

In the equation for Cohen's  $d_s$  (see below), the numerator is the difference between the appropriate means of two groups of observations. The denominator is the pooled standard deviation. These differences are squared to prevent the positive and negative values cancelling each other out. Then, they are summed. This is then divided by the number of observations minus one (referred to as Bessel's correction) for bias in the estimation of the population variance, and then the square root of this is taken.

$$\text{Cohen's } d_s = \frac{\bar{X}_1 - \bar{X}_2}{\sqrt{\frac{(n_1 - 1)SD_1^2 + (n_2 - 1)SD_2^2}{n_1 + n_2 - 2}}}$$

Regarding these effect sizes  $d_s$ , the following describes the magnitude for the values according to Sawilowsky<sup>29</sup>; 0.01 > very small effect; 0.20 > small effect; 0.50 > medium effect; 0.80 > large effect; 1.20 > very large effect, and finally 2.00 > a huge effect size.

### Cubic-spline interpolation

Subsequently, we assessed the importance of pre-processing techniques on the results obtained through chaotic global algorithms. Again, we compare the chaotic global values for CFP1–CFP7. Time series constructed from the RR-interval tachograms are not equidistantly sampled. This has to be justified before frequency-domain analysis.

Primarily, we can decide to assume equidistant sampling<sup>30</sup> and compute the power spectrum directly from the tachogram of RR-intervals. This is the technique widely adopted up-til-now by previous studies on chaotic globals with obese children,<sup>31</sup> Attention Deficit Hyperactivity Disorder (ADHD),<sup>23</sup> type 1 diabetes mellitus,<sup>32</sup> and flexible-pole physical therapy shoulder rehabilitation<sup>33</sup> among others. The RR-intervals are, therefore, a function of the

**Table 1.** Sex, mean values followed by their respective standard deviations of age, mass, and body mass index.

Variable	Obese	Control
Sex (female/male)	20/23	21/22
Age (years)	20.45 ± 1.57	20.70 ± 1.39
Mass (kg)	102.30 ± 20.82	62.89 ± 10.47
Body mass index (kg/m <sup>2</sup> )	34.67 ± 3.87	21.91 ± 1.86

beat number. Yet, this could cause a distortion in the spectrum<sup>34</sup> and the spectrum must be considered a function of cycles per beat rather than of frequency.<sup>35</sup> Incidentally, a Lomb power spectrum<sup>36</sup> could also be used as an alternative power spectrum specifically for unequally spaced data. Nevertheless, here it is not relevant so not discussed further.

A completely different approach here is to enforce a cubic spline interpolation<sup>37</sup> to convert the non-equidistantly sampled RR-tachogram into an equidistantly sampled time series.<sup>38</sup> So, we performed a cubic spline interpolation on the RR-interval tachogram. We accomplished this at the levels 1–13 Hz. This covers most relevant scenarios in heart rate variability analysis. Kubios HRV<sup>®39</sup> software offers a default option of 4 Hz. It is important to grasp that the interpolation frequency will affect the number of data points in the time series. A frequency of 4 Hz, for example, will elevate the number of RR-intervals from 125 (1 Hz) to 500 (4 Hz).

Following the cubic spline interpolation, the chaotic global algorithms parameters are fixed. Throughout the enforcement of the cubic spline interpolations, the Multi-Taper Method default parameters were set to the following: Thomson's setting to "adaptive," sampling frequency to 1 Hz, Fast Fourier Transform (FFT) length of 256 and the discrete prolate spheroidal sequence to 3.

## Results

Table 1 presents data regarding (sex, age, mass, and body mass index).

Statistics illustrate that there is a wide variation in both the mean values and standard deviation for both non-obese and obese youths. Here, we are assessing ultra-short time series (125 RR-intervals). Previously, in studies of obese youths,<sup>7,8</sup> 1000 RR-intervals were evaluated. The chaotic global algorithms in this study compute a significant statistical result ( $p < 0.01$ ,  $< 1\%$ ) for three of the seven combinations (see Table 2 and Fig 1). These are combinations CFP1, CFP3 and CFP6 as with the former study but, here with an 8 times shorter time series. In all three cases, there is a significant increase in chaotic response when comparing non-obese to obese youth patients.

The non-parametric Kruskal–Wallis test of significance was calculated. This was since the distributions were revealed to be non-normal in the majority of cases, determined by the Anderson–Darling<sup>40</sup> and Lilliefors<sup>41</sup> statistical tests. Effect sizes by Cohen's  $d_s$ <sup>26–28</sup> were also calculated as there was a wider range of values.

With regard to the multivariate statistical analysis by principal component analysis (see Fig 2). Only the first two components need be considered by reason of a moderately steep scree plot. The cumulative influence as a percentage is 66.8% for Principal Component 1 (PC1) and 99.8% for the cumulative total of the PC1 and Principal Component 2 (PC2). PC2 has an individual

**Table 2.** Mean values and standard deviations for the chaotic forward parameters CFP1 to CFP7 which are non-dimensional values; for the non-obese and obese youth patients with 125 RR-intervals. Kruskal–Wallis test of significance was computed as distributions were mainly non-normal by Anderson–Darling and Lilliefors tests of normality. Cohen's  $d_s$  effects sizes were also calculated where a negative value signifies an increase in chaotic response from non-obese to obese youth and a positive value the opposite response.

Chaotic global CFPx	Mean ± standard deviation, normal (n=43)	Mean ± standard deviation, youth obese (n=43)	Kruskal–Wallis (p-value)	Cohen's $d_s$ effect sizes
CFP1	0.8080 ± 0.1154	0.8726 ± 0.1225	0.001	–0.542
CFP2	0.5995 ± 0.1036	0.5828 ± 0.0795	0.723	0.181
CFP3	0.6960 ± 0.1080	0.7613 ± 0.0957	<0.001	–0.640
CFP4	0.6536 ± 0.1994	0.7611 ± 0.1986	0.015	–0.540
CFP5	0.3819 ± 0.1574	0.4106 ± 0.1394	0.364	–0.193
CFP6	0.5254 ± 0.1428	0.6382 ± 0.1532	0.001	–0.761
CFP7	0.3965 ± 0.2090	0.3571 ± 0.1775	0.374	0.203

influence of 33.0%. From Figure 2, it is evident that metrics CFP3 and CFP1 are the most influential on the basis of the first two components. This is to be expected as CFP1 is usually the most statistically robust and CFP3 the most statistically significant.

The effect of cubic spline interpolation between 1 and 13 Hz increasing the length of the time series by interpolation (rather than by recording longer time series in the laboratory) is negligible. This is for CFP1, CFP3 and CFP6 via both Kruskal–Wallis and effect sizes by Cohen's  $d_s$  test of significances as illustrated in Table 3. It is apparent that the Cohen's  $d_s$  are most significant for CFP6 ( $d_s \approx -0.76$ ) then CFP3 ( $d_s \approx -0.64$ ), and least so for CFP1 ( $d_s \approx -0.54$ ). The negative sign indicates an increase from non-obese to obese youths. The Kruskal–Wallis tests indicate significances of  $p < 0.01$  for all, hence, the use of Cohen's  $d_s$  which has a wider range of values and so is more useful to discriminate between the interpolations.

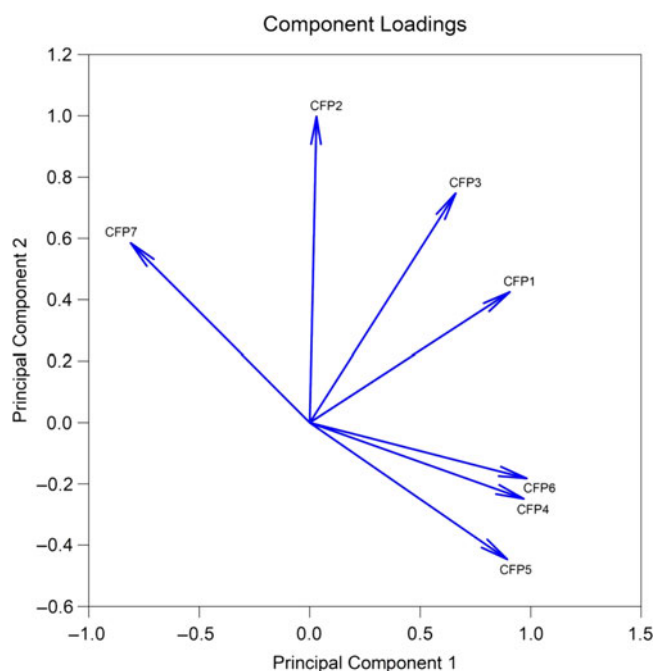
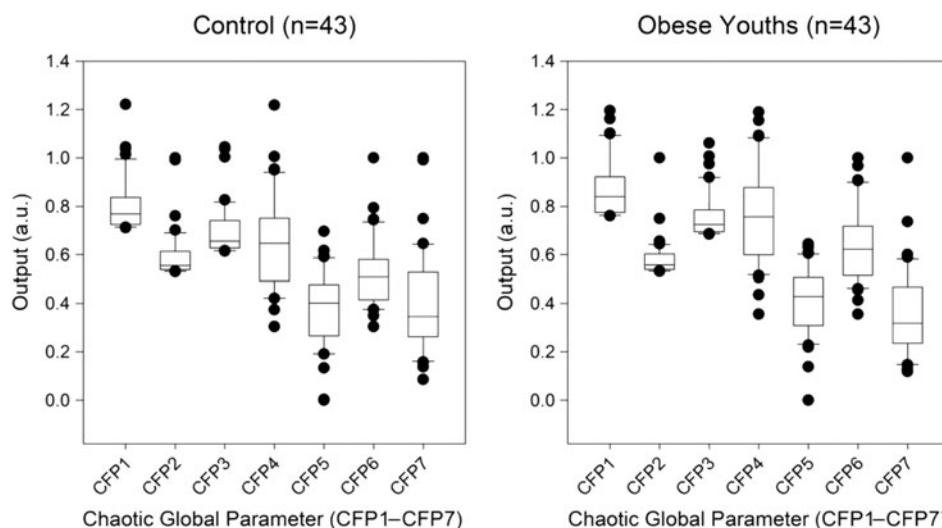
## Discussion

It is unclear why different algorithms behave in alternative ways for heart rate autonomic control. Consequently, our study aimed to assess a new approach to detect autonomic dysfunction in obese youths based on the non-linear dynamics from the non-periodic RR-intervals oscillations. As a chief outcome, chaotic global techniques applied for heart rate variability analysis were able to identify cardiac autonomic dysfunction in a sample of obese youths and using an ultra-short time series of 125 RR-intervals.

Heart rate variability has received much attention by reason of its simple workability. Data can be collected by a one-channel electrocardiographic or a pulse watch. Then, these RR-intervals can be processed by Kubios HRV<sup>®</sup> software.<sup>39</sup> Previously, Task Force in 1996 published directives in order to regulate heart rate variability analysis using linear methods in the time and frequency domains.<sup>38</sup>

Non-linear analysis of heart rate variability was specified to provide information about the scaling, quality, and correlation properties of the time series. There are countless non-linear techniques; some based on Approximate, Sample, Shannon, Renyi and Tsallis Entropies<sup>42</sup> or, Higuchi and Katz's fractal dimensions<sup>43</sup> and

**Figure 1.** The boxplots for CFP1 to CFP7 aimed at non-obese and the obese youth patients (n=43) with 125 RR-intervals. The point closest to the zero is the minimum and the point farthest away is the maximum. The point next closest to the zero is the 5th percentile and the point next farthest away is the 95th percentile. The boundary of the box closest to zero indicates the 25th percentile, a line within the box marks the median (not the mean), and the boundary of the box farthest from zero indicates the 75th percentile. The difference between these points is the inter-quartile range. Whiskers (or error bars) above and below the box indicate the 90th and 10th percentiles, respectively.



**Figure 2.** The plot illustrates the component loadings CFP1 to CFP7 for the 125 RR-intervals of 43 obese youth patients' described above with a cubic spline interpolation of 1 Hz. The Chaotic Forward Parameter values are deduced by using the Multi-Taper Method spectra throughout. The properties of the Multi-Taper Method spectra are as follows: Sampling frequency 1 Hz, discrete prolate spheroidal sequence of 3, FFT length of 256 and Thomson's non-linear combination at "adaptive." CFP1 and CFP3 perform best when assessed by principal component analysis; the most influential components.

specifically the novel chaotic global metrics<sup>17</sup> investigated in this study. Linear methods were intended to compute heart rate variability's extent. Complex algorithms to assess the level of chaotic response of heart rate variability are suggested to detect autonomic changes that linear methods are unable to identify.<sup>44</sup> This is the key advantage of the non-linear techniques.

Globally, chaotic methods have hitherto been applied to RR-intervals in obese children<sup>31</sup> or obese youths<sup>7</sup> and, malnourished

children.<sup>45</sup> There have, thus far, been no studies enforcing the cubic spline interpolation on chaotic global methods. Few studies have assessed ultra-short time series of 125 RR-intervals with any metric. The previous studies<sup>7,23</sup> have assessed time series 8 times lengthier. Historically, metrics assessing data required 24-hour Holter electrocardiographic recordings<sup>46</sup> to make considerations. This was severely reduced when the spectrally involved chaotic globals were introduced.<sup>17</sup> Until now, even chaotic globals had not been tested on 125 RR-intervals. Ultimately, the results are favorable with three metrics (CFP1, CFP3 and CFP6) all discriminated from the controls at the level  $p < 0.01$  (or,  $< 1\%$ ) on ultra-short time series.

Some limitations from our study need highlighting. We evaluated a small sample, yet statistical analysis provided significance. The sample was comprised of only Brazilian patients. Thus, we should be cautious when interpreting such data in countries from different continents. We did not obtain information regarding body fat percentage, lean mass, inflammatory markers, and oxidative stress. Further studies are encouraged to evaluate the correlations between the mentioned variables and chaotic global analysis. Different autonomic approaches, for example, electroneuromyography, baroreflex function, and skin response were not investigated, as our emphasis was chaotic global analysis applied to RR-intervals – a relatively low-cost technique.

Our study presents important findings for clinical practice and procedures. ICUs and physicians are interested in predicting the risk for physiological complications. Comprehension of biological signals through non-linear analysis of heart rate variability is a significant issue for an appropriate program of care. We revealed that chaotic globals applied to ultra-short time series of RR-intervals are sensitive to differentiate autonomic impairment of obese youths from non-obese youth patients. Yet, cubic spline interpolations have only trivial effects. Therefore, we can accept that cubic spline interpolation is unnecessary, and shorter than usual time series are adequate to make decisions about cardiac autonomic dysfunction in obese youths. By using shorter time series computations, they are less processor intensive and can be calculated faster which is advantageous in an ICU setting where decisions need to be made quickly. Nonetheless, it is important to realise though this may not be the case with other experimental groups which must be assessed individually on their merits alone.

**Table 3.** CFP1, CFP3 and CFP6 (non-dimensional values) and their test of significances by Kruskal–Wallis and effect sizes Cohen's  $d_s$  for the non-obese compared to obese youth patients. Discrete prolate spheroidal sequence is set to 3 and length of time series starts at 125 RR-intervals for an interpolation rate of 1 Hz. So, this increases to 250 RR-intervals with an interpolation rate of 2 Hz and so on. To calculate the Multi-Taper Method, settings are fixed as follows: a sampling frequency of 1 Hz, 256 for FFT length and Thomson's "adaptive" non-linear combination method.

Interpolation rate (Hz)	Kruskal–Wallis test			Effect sizes by Cohen's $d_s$		
	CFP1	CFP3	CFP6	CFP1	CFP3	CFP6
1	0.001	<0.001	0.001	−0.543	−0.640	−0.761
2	0.001	<0.001	0.001	−0.545	−0.639	−0.760
3	0.001	<0.001	0.001	−0.546	−0.640	−0.761
4	0.001	<0.001	0.001	−0.547	−0.639	−0.760
5	0.001	<0.001	0.001	−0.546	−0.640	−0.760
6	0.001	<0.001	0.001	−0.547	−0.640	−0.760
7	0.001	<0.001	0.001	−0.547	−0.639	−0.759
8	0.001	<0.001	0.001	−0.549	−0.639	−0.760
9	0.001	<0.001	0.001	−0.548	−0.640	−0.760
10	0.001	<0.001	0.001	−0.548	−0.639	−0.760
11	0.001	<0.001	0.001	−0.548	−0.640	−0.760
12	0.001	<0.001	0.001	−0.548	−0.640	−0.760
13	0.001	<0.001	0.001	−0.548	−0.639	−0.760

## Conclusion

The three chaotic global techniques (CFP1, CFP3, and CFP6) applied to an ultra-short time series of 125 RR-intervals robustly detected heart rate variability deviations in obese youth patients. Extensive interpolation of time series made statistically insignificant effects by two statistical tests, and so was unnecessary. Yet, these three techniques were able to identify autonomic dysfunction in obese youth patients and accordingly discriminate between these two groups.

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**Conflicts of Interest.** None.

**Ethical Standards.** All procedures performed in studies involving human patients were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. All volunteers signed a consent letter and was informed of the procedures and objectives of the study. The study's procedures were all approved by the Research Ethics Committee of Sao Paulo State University, UNESP (Number Protocol No. 11/2011).

## References

- Kleiger RE, Miller JP, Bigger JT, Moss AJ. Decreased heart rate variability and its association with increased mortality after acute myocardial infarction. *Am J Card* 1987; 59: 256–262.
- Campos LA, Pereira VL Jr., Muralikrishna A, Albarwani S, Brás S, Gouveia S. Mathematical biomarkers for the autonomic regulation of cardiovascular system. *Front Physiol* 2013; 4: 279.
- Wiertel-Krawczuk A, Hirschfeld AS, Huber J, Wojtyasiak M, Szymankiewicz-Szukala A. Sympathetic skin response following single and combined sound and electrical stimuli in young healthy subjects. *J Med Sc* 2016; 85: 106–113.
- Baum P, Petroff D, Classen J, Kiess W, Bluher S. Dysfunction of autonomic nervous system in childhood obesity: a cross-sectional study. *PLoS One* 2013; 8: e54546.
- Goldberger AL. Cardiac chaos. *Science* 1989; 243: 1419.
- Goldberger AL, West BJ. Chaos and order in the human body. *MD Comput* 1992; 9:25–34.
- Vanderlei FM, Vanderlei LCM, Garner DM. Heart rate dynamics by novel chaotic globals to HRV in obese youths. *J Hum Growth Develop* 2015; 25: 82–88.
- Garner DM, Vanderlei FM, Vanderlei LCM. Complex measurements of heart rate variability in obese youths: distinguishing autonomic dysfunction. *J Hum Growth Develop* 2018; 28: 298–306.
- McKinley S, Levine M. Cubic spline interpolation. *Coll Red* 1998; 45: 1049–1060.
- Mackey MC, Milton JG. Dynamical diseases. *Ann N Y Acad Sci* 1987; 504: 16–32.
- Bélair J, Glass L, An der Heiden U, Milton J. Dynamical disease: identification, temporal aspects and treatment strategies of human illness. *Chaos* 1995; 5: 1–7.
- Seiver A, Daane S, Kim R. Regular low frequency cardiac output oscillations observed in critically ill surgical patients. *Complexity* 1997; 2: 51–55.
- Kawaguchi M, Takamatsu I, Kazama T. Rocuronium dose-dependently suppresses the spectral entropy response to tracheal intubation during propofol anaesthesia. *Br J Anaesth* 2009; 102: 667–672.
- Alvarez D, Hornero R, Marcos J, Del Campo F, Lopez M. Spectral analysis of electroencephalogram and oximetric signals in obstructive sleep apnea diagnosis. *Conf Proc IEEE Eng Med Biol Soc* 2009; 2009: 400–403.
- Bokov P, Fiamma MN, Chevalier-Bidaud B, et al. Increased ventilatory variability and complexity in patients with hyperventilation disorder. *J Appl Physiol* 2016; 120: 1165–1172.
- Grogono JC, Butler C, Izadi H, Moosavi SH. Inhaled furosemide for relief of air hunger versus sense of breathing effort: a randomized controlled trial. *Respir Res* 2018; 19: 181.
- Garner DM, Ling, BWK. Measuring and locating zones of chaos and irregularity. *J Syst Sci Complex* 2014; 27: 494–506.
- Ghil M. The SSA-MTM toolkit: applications to analysis and prediction of time series. *Appl Soft Comp* 1997; 3165: 216–230.
- Garner DM, De Souza NM, Vanderlei LCM. Risk assessment of diabetes mellitus by chaotic globals to heart rate variability via six power spectra. *Rom J Diabetes Nut Met Dis* 2017; 24: 227–236.
- Slepian S. Prolate spheroidal wave functions, Fourier analysis and uncertainty, V, The discrete case. *Bell Syst Tech J* 1978; 57: 1371–1430.
- Shannon CE. A mathematical theory of communication. *Bell Syst Tech J* 1948; 27: 379–423.
- Peng CK, Havlin S, Stanley HE, Goldberger AL. Quantification of scaling exponents and crossover phenomena in nonstationary heartbeat time series. *Chaos* 1995; 5: 82–87.
- Wajnsztein R, De Carvalho TD, Garner DM, et al. Heart rate variability analysis by chaotic global techniques in children with attention deficit hyperactivity disorder. *Complexity* 2016; 21: 412–419.
- Jolliffe IT. *Principal Component Analysis*, Springer Series in Statistics, vol. 2nd edn. Springer, New York, 2002.
- Manly BF. *Multivariate Statistical Methods: A Primer*. CRC Press, Boca Raton, 2004.
- Sullivan GM, Feinn R. Using effect size—or why the P value is not enough. *J Grad Med Ed* 2012; 4: 279–282.
- Coe R. It's the Effect Size, Stupid: what effect size is and why it is important. *British Educational Research Association annual conference*. 2002; 1–18. Accessed at: <https://www.cem.org/attachments/ebe/ESguide.pdf>
- Lakens D. Calculating and reporting effect sizes to facilitate cumulative science: a practical primer for t-tests and ANOVAs. *Front Psychol* 2013; 4: 863.
- Sawilowsky SS. New Effect Size Rules of Thumb 2009; 8: 597–599.
- Baselli G, Cerutti S, Civardi S, et al. Heart rate variability signal processing: a quantitative approach as an aid to diagnosis in cardiovascular pathologies. *Int J Biom Comp* 1987; 20: 51–70.

31. Vanderlei FM, Vanderlei LC, Garner DM. Chaotic global parameters correlation with heart rate variability in obese children. *J Hum Growth Develop* 2014; 24: 24–30.
32. Souza NM, Vanderlei LC, Garner DM. Risk evaluation of diabetes mellitus by relation of chaotic globals to HRV. *Complexity*. 2015; 20: 84–92.
33. Antonio AMS, Garner DM, Cardoso MA, et al. Behaviour of globally chaotic parameters of heart rate variability following a protocol of exercise with flexible pole. *Russ J Cardiol* 2015; 4: 24–28.
34. Mateo J, Laguna P. Improved heart rate variability signal analysis from the beat occurrence times according to the IPPFM model. *IEEE Trans Biomed Eng* 2000; 47: 985–996.
35. DeBoer RW, Karemaker JM, Strackee J. Comparing spectra of a series of point events particularly for heart rate variability data. *IEEE Trans Biomed Eng* 1984; 31(4): 384–387.
36. Lomb NR. Least-squares frequency analysis of unequally spaced data. *Astrophys Space Sci* 1976; 39: 447–462.
37. Kreyszig E. *Advanced Engineering Mathematics*. Wiley, Hoboken, 2011.
38. Camm AJ, Malik M, Bigger JT, et al. Heart rate variability: standards of measurement, physiological interpretation and clinical use. Task Force of the European Society of Cardiology and the North American Society of Pacing and Electrophysiology. *Circulation* 1996; 93: 1043–1065.
39. Tarvainen MP, Niskanen JP, Lipponen JA, Ranta-Aho PO, Karjalainen PA. Kubios HRV—heart rate variability analysis software. *Comp Meth Prog Biomed* 2014; 113: 210–220.
40. Anderson TW, Darling DA. A test of goodness of fit. *J Am Stat Assoc* 1954; 49: 765–769.
41. Razali NM, Wah YB. Power comparisons of Shapiro-Wilk, Kolmogorov-Smirnov, Lilliefors and Anderson-Darling tests. *J Stat Mod Anal* 2011; 2: 21–33.
42. Vanderlei F, Vanderlei LCM, de Abreu LC, Garner DM. Entropic analysis of HRV in obese children. *Int Arch Med* 2015; 8(200): 1–9.
43. Garner DM, Souza NM, Vanderlei LCM. Heart rate variability analysis: Higuchi and Katz's fractal dimensions in subjects with type 1 diabetes mellitus. *Rom J Diabetes Nut Met Dis* 2018; 25: 289.
44. Sassi R, Cerutti S, Lombardi F, et al. Advances in heart rate variability signal analysis: joint position statement by the e-Cardiology ESC Working Group and the European Heart Rhythm Association co-endorsed by the Asia Pacific Heart Rhythm Society. *Ep Europace* 2015; 17: 1341–1353.
45. Barreto GS, Vanderlei FM, Vanderlei LCM, Garner DM. Risk appraisal by novel chaotic globals to HRV in subjects with malnutrition. *J Hum Growth Develop* 2014; 24: 243–248.
46. Van Leeuwen P, Bettermann H, An der HU, Kummell HC. Circadian aspects of apparent correlation dimension in human heart rate dynamics. *Am J Physiol* 1995; 269: H130–H134.