

Multi-lag Tone - Entropy in Neonatal Stress

Matej Šapina, Matej; Karmakar, Chandan Kumar; Kramarić, Karolina; Garcin, Matthieu; Adelson, P. David; Milas, Krešimir; Pirić, Marko; Brdarić, Dario; Yearwood, John

Source / Izvornik: **Journal of the Royal Society Interface, 2018, 15, 1 - 8**

Journal article, Published version

Rad u časopisu, Objavljena verzija rada (izdavačev PDF)

<https://doi.org/10.1098/rsif.2018.0420>

Permanent link / Trajna poveznica: <https://urn.nsk.hr/urn:nbn:hr:239:488924>

Rights / Prava: [Attribution 4.0 International](#)/[Imenovanje 4.0 međunarodna](#)

Download date / Datum preuzimanja: **2024-12-26**



Repository / Repozitorij:

[Repository UHC Osijek - Repository University Hospital Centre Osijek](#)



Research



Cite this article: Šapina M, Karmakar CK, Kramarić K, Garcin M, Adelson PD, Milas K, Pirić M, Brdarić D, Yearwood J. 2018 Multi-lag tone–entropy in neonatal stress. *J. R. Soc. Interface* **15**: 20180420.
<http://dx.doi.org/10.1098/rsif.2018.0420>

Received: 7 June 2018
Accepted: 29 August 2018

Subject Category:
Life Sciences—Physics interface

Subject Areas:
bioengineering, biomedical engineering, medical physics

Keywords:
heart rate variability, newborn, stress, tone–entropy, autonomic nervous system

Author for correspondence:
Matej Šapina
e-mail: sapina.matej@yahoo.com

†These authors contributed equally to this study.

Multi-lag tone – entropy in neonatal stress

Matej Šapina^{1,2,3,†}, Chandan Kumar Karmakar^{4,5,†}, Karolina Kramarić^{1,2,3},
Matthieu Garcin⁶, P. David Adelson⁷, Krešimir Milas^{1,2}, Marko Pirić²,
Dario Brdarić^{3,8} and John Yearwood⁴

¹University hospital Osijek, Pediatric Clinic, J. Huttlera 4, 31000 Osijek, Croatia

²Medical faculty Osijek, Osijek, Cara Hadrijana 10E, 31000 Osijek, Croatia

³Faculty of Dental medicine and Health, Crkvena 21, 31000 Osijek, Croatia

⁴School of Information Technology, Deakin University, Geelong, Australia

⁵Department of Electrical and Electronic Engineering, University of Melbourne, Melbourne, Australia

⁶LabEx ReFi, 106 boulevard de l'Hôpital, 75013 Paris, France

⁷Barrow Neurological Institute at Phoenix Children's Hospital, Phoenix, AZ, USA

⁸Institute of Public Health for the Osijek-Baranya County, Drinska 8, 31000 Osijek, Croatia

MŠ, 0000-0002-4376-708X

Heart rate variability (HRV) has been analysed using linear and nonlinear methods. In the framework of a controlled neonatal stress model, we applied tone–entropy (T–E) analysis at multiple lags to understand the influence of external stressors on healthy term neonates. Forty term neonates were included in the study. HRV was analysed using multi-lag T–E at two resting and two stress phases (heel stimulation and a heel stick blood drawing phase). Higher mean entropy values and lower mean tone values when stressed showed a reduction in randomness with increased sympathetic and reduced parasympathetic activity. A ROC analysis was used to estimate the diagnostic performances of tone and entropy and combining both features. Comparing the resting and simulation phase separately, the performance of tone outperformed entropy, but combining the two in a quadratic linear regression model, neonates in resting as compared to stress phases could be distinguished with high accuracy. This raises the possibility that when applied across short time segments, multi-lag T–E becomes an additional tool for more objective assessment of neonatal stress.

1. Introduction

Neonatal stress is a special field of research and clinical practice that can be considered from a wide spectrum of perspectives [1–3]. Endocrinologically, during a stress phase, there is a large excretion of hormones generated to cope with a potentially life-threatening event, further activating a part of the autonomic nervous system (ANS), and producing significant changes in every body system—basically changing the body's homeostasis [4–6]. These described changes are a result of the activation of the sympathetic branch of the ANS, which can result in an increase in the heart and respiratory rate, blood pressure, causing an increase of stress hormone levels (i.e. adrenaline, noradrenaline, cortisol, etc.). By contrast, the parasympathetic branch predominates, when there is no need for an acute stress response. The activation of the parasympathetic branch of the ANS decreases the heart and respiratory rate, as well as blood pressure, increases salivation, urination, digestion, defecation, all of which should be delayed in acute stress situations [7,8]. Owing to the constant interaction with various stimuli, the perturbations in the predominance of one of the two branches of the ANS constantly occurs, trying to maintain homeostasis. However, imbalances, such as prolonged activation of the sympathetic or inhibition of the parasympathetic branch, are considered risk factors in various diseases, which can ultimately lead to death (i.e. ventricular tachyarrhythmias and sudden cardiac death) [9,10].

Clinically, neonates and especially those born prematurely can sometimes present with vague clinical signs and symptoms of disease [11–14]. As children physiologically differ from adults, stressors that do not affect adult humans

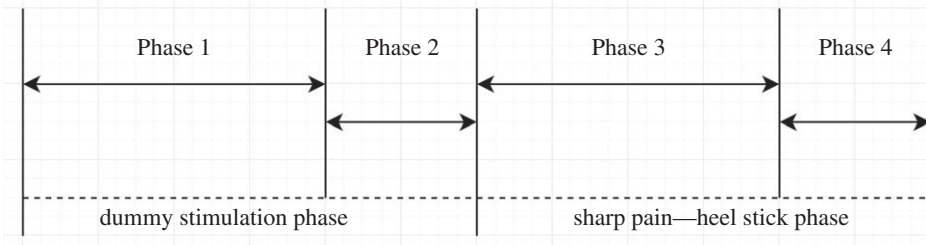


Figure 1. The sequence of phases used in the study protocol. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 3, second baseline phase; Phase 4, heel stick blood drawing phase.

might have a significant impact on the very young, altering their long-term development [15,16]. In addition to environmental stressors naturally characteristic of the neonatal intensive care unit (NICU), (i.e. noise, temperature, etc.) if the neonate is further challenged with disease or early gestational age at birth, they experience more stress from various invasive medical procedures they undergo [17].

Heart rate variability (HRV) is the variability of the duration between successive cardiac cycles that originate in the sinus node [18]. HRV has broad application in basic ANS science and clinical use via the application of different mathematical and statistical methods: time and spectral analysis, measures of entropy, Hurst exponents, indices derived from chaos theory such as the correlation dimension, Poincaré plots, recurrence plots or Lyapunov exponents [19–22].

Among the various HRV parameters, tone–entropy (T–E) showed promising results in identifying patients with ANS dysfunction [23]. T–E uses successive RR intervals with the implicit assumption that the current heartbeat is influenced by the immediately preceding beat. However, it has been reported that each heartbeat influences not only the beat immediately following it, but also up to 6–10 beats downstream, possibly as a consequence of respiratory sinus arrhythmia and baroreflex influence [24]. In addition, several researchers have shown that the delayed signal can be used for measuring the influence on the succeeding beats [25] and the delay is associated with the disease processes [24]. Thus, successive RR interval duplets will likely underestimate the role of the autocovariance function of RR intervals, i.e. the ability of heart beats to influence a train of succeeding beats. The strong correlation between successive beats (in case of lag 1 T–E) also masks the nonlinearity from the measurement [23,26]. Thus, use of multi-lag analysis can enable the autocovariance function to capture a nonlinear aspect of HRV. In a previous study [19], it has been reported that multi-lag T–E analysis can overcome the limitations of the single-lag T–E analysis in HRV studies. They showed that multi-lag T–E analysis better stratified the risk of autonomic neuropathy in Type 2 diabetes mellitus than single-lag T–E [19]. In addition, a rationale for the use of T–E relies on the evidence that T–E is not influenced by the period of data collection or the baseline heart rate (HR) [27].

2. Subjects and methods

2.1. The study protocol

Forty healthy term neonates (21 females and 19 males, birth weight 3542.05 ± 339.09 g), all of who were born through

vaginal delivery, with an APGAR score $>9/9$ at 1 and 5 min, without perinatal risk factors, breastfed or additionally formula fed if not enough breastmilk was supplied, and not previously experiencing any external stress stimuli, were included in this study. The neonates were chosen by simple random sampling over a time period from 30 September 2016 to 10 February 2017, in the paediatrics clinic’s maternity ward of the Osijek University Clinic. Among the participants, only two female neonates were twins, while the rest of the participants were singletons. The average gravidity and parity of the participant’s mothers were 1.66 ± 0.92 . All mothers were euthyroid, one participant’s mother was diagnosed with gestational diabetes, but did not require treatment with insulin therapy, and none of the mothers had prior recorded miscarriages. The experimental paradigm was performed at the recommended chronological age for routine metabolic screening, 72 h of age, just prior discharge [28,29]. None of the neonates included in this study underwent any other procedural pain besides the national metabolic screening heel stick blood draw. As we have described previously [30], the protocol is divided into three parts: (a) dummy stimulation, (b) the sharp pain—heel stick, (c) the treatment; only the first two parts were included in the study (figure 1).

Part (a) consisted of two phases: the first, the baseline phase (no stimulation), lasting 10 min (further named Phase 1), after which intermittent pressing of the neonate’s heel was done, mimicking a heel stick blood drawing procedure without actual blood sampling (Phase 2). Phase 2 lasted 90 s, which is the average time for the nurse to perform the blood drawing procedure. The end of Phase 2 is the starting point of Part (b). Part (b) also consisted of two phases: Phase 3, the second baseline (no stimulation), again lasting 10 min, followed by the actual heel stick blood sampling procedure (Phase 4).

A lightweight, high-resolution device, with a high sampling rate (1024 Hz) was used (Firstbeat Bodyguard 2, Firstbeat Technologies Ltd, Jyvaskyla, Finland) during the entire procedure. After visual inspection and artefact removal, the data were analysed. To reduce movement artefacts and ensure quality of the recordings, the neonates were positioned supine after breast- or formula feeding in a quiet room.

2.2. Multi-lag T–E analysis of HRV signal

The RR interval is defined as the time difference between two consecutive R peaks of the electrocardiogram signal and the RR intervals in time series (RR_i) are defined as

$$RR = (RR_1, RR_2, \dots, RR_N),$$

Table 1. Mean \pm s.d. of tone values across the different phases of the protocol with varying lags. P -values were calculated using one-way ANOVA, and represent significance of differences among tone values obtained at different phases. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 3, second baseline phase; Phase 4, heel stick blood drawing phase.

lag	Phase 1	Phase 2	Phase 3	Phase 4	P^*
1	-0.09 ± 0.10	0.04 ± 0.09	-0.09 ± 0.11	-0.02 ± 0.13	<0.001
2	-0.12 ± 0.10	0.11 ± 0.18	-0.12 ± 0.11	0.00 ± 0.19	<0.001
3	-0.17 ± 0.12	0.19 ± 0.27	-0.17 ± 0.15	0.03 ± 0.26	<0.001
4	-0.20 ± 0.13	0.26 ± 0.37	-0.21 ± 0.17	0.05 ± 0.34	<0.001
5	-0.24 ± 0.15	0.34 ± 0.47	-0.24 ± 0.20	0.08 ± 0.42	<0.001
6	-0.27 ± 0.16	0.42 ± 0.56	-0.27 ± 0.22	0.11 ± 0.51	<0.001
7	-0.31 ± 0.17	0.49 ± 0.66	-0.30 ± 0.24	0.14 ± 0.60	<0.001
8	-0.33 ± 0.18	0.57 ± 0.76	-0.33 ± 0.26	0.17 ± 0.70	<0.001
9	-0.35 ± 0.19	0.64 ± 0.86	-0.35 ± 0.28	0.21 ± 0.81	<0.001
10	-0.38 ± 0.20	0.72 ± 0.96	-0.36 ± 0.29	0.24 ± 0.91	<0.001

where N is the number of RR_i . HR acceleration and inhibition can be determined from the difference of consecutive RR_i . If RR_{i+1} becomes shorter than RR_i , then it is an acceleration of HR. Therefore, acceleration of the heart is expressed as a plus difference and inhibition as a minus difference of RR_i . However, to reduce the impact of HR variation over a wide range of time and different subjects, normalized variation in RR_i is preferred to monitor the variability. This normalized variation is measured by percentile change (percentage index (PI)) and defined as:

$$PI_i^m = \frac{RR_i - RR_{i+m}}{RR_i} \times 100, \quad (2.1)$$

where m is an integer and represents the lag used for measuring the PI. The detailed methodology of multi-lag T–E analysis has been described in a previous report [23]. The *Tone* at lag m ($Tone^m$) is defined as a first-order moment (arithmetic average) of this PI^m time series as

$$Tone^m = \frac{1}{N-m} \sum_{i=1}^{N-m} PI_i^m. \quad (2.2)$$

The *Entropy* at lag m is defined from the probability distribution of PI^m by using Shannon's formula [31]:

$$Entropy^m = - \sum_{i=1}^n p(i) \log_2 p(i), \quad (2.3)$$

where $p(i)$ is a probability that PI_n^m has a value in the range $i \leq PI_n^m < i+1$, i is an integer. The $Entropy^m$ evaluates total acceleration–inhibition activities, or total heart period variations, in a familiar unit of bit.

2.3. Statistical analysis

The data were analysed with the Matlab R2007a software. The Kolmogorov–Smirnov test was used to test the normality of distributions. The data are descriptively presented with means and standard deviations. A repeated measures ANOVA, followed by a pairwise post hoc test, was applied to assess the mean differences of tone and entropy across the different phases. An ROC curve analysis was used to test the diagnostic properties of the T–E comparing the stress phases (Phase 2 and 4) to the first baseline (Phase 1). The results of the ROC curve analysis are presented as the total area under

the ROC curve (AUC), which represents the performance of the model. AUC values enable comparison of the performance of different models, and it is generally accepted that models with higher AUC values have a better overall performance of correctly classifying diseased (in our case stress phase) from non-diseased (resting phase) subjects. P -values less than 0.05 were considered statistically significant.

3. Results

There were statistically significant differences ($p < 0.001$) when comparing both the tone and entropy values across the four phases during the first 10 lags (tables 1 and 2). Over the varying lags, at the resting phases (Phase 1 and 3), a steady linear decrease of tone values was observed ranging from -0.09 ± 0.10 in Phase 1, and -0.09 ± 0.11 , at lag 1, up to -0.38 ± 0.20 in Phase 1, and -0.36 ± 0.29 (Phase 3) (table 1). On the contrary, at both stress phases (Phase 2 and Phase 4), a steady linear increase in the tone values is observed. The tone values at Phase 2 were always positive, being the lowest at lag 1 (0.04 ± 0.09), and highest at lag 10 (0.72 ± 0.96), while a single negative tone value was observed at lag 1 of Phase 4 (-0.02 ± 0.13). At every lag the tone values of Phase 4, were lower than in Phase 2, being highest at Phase 4 (0.24 ± 0.91).

Contrary to the tone values which decrease in the stress phases (table 1), a steady increase of entropy is observed at all phases (table 2). At each lag, the entropy values were observed to be lower at stress (Phase 2 and 4) than at rest (Phase 1 and 3). The lowest entropy at all phases was observed at lag 1 (Phase 1: 3.63 ± 0.63 , Phase 2: 3.11 ± 0.59 , Phase 3: 3.58 ± 0.74 , Phase 4: 4.28 ± 0.60), while the highest values were observed at lag 10 (Phase 1: 4.87 ± 0.46 , Phase 2: 4.26 ± 0.59 , Phase 3: 4.26 ± 0.59 , Phase 4: 4.28 ± 0.60).

The post hoc p -values comparing both tone and entropy across the phases are presented in table 3. When comparing both tone and entropy values in the resting phases (Phase 1 and Phase 3), no significant differences were observed. Tone significantly differed between Phase 1 and 2 (p -values: from lag 1 to lag 10 – <0.001), and Phase 1 and 4 at all lags (p -values: lag 1 – 0.015, lag 2 – 0.001, from lag 3 to lag 10 – <0.001), as well as between Phase 3 and Phase

Table 2. Mean \pm s.d. of entropy values across the different phases of the protocol with varying lags. P -values were calculated using one-way ANOVA, and represent significance of differences among entropy values obtained at different phases. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 3, second baseline phase; Phase 4, heel stick blood drawing phase.

lag	feature	Phase 1	Phase 2	Phase 3	Phase 4	P^*
1	entropy	3.63 \pm 0.63	3.11 \pm 0.59	3.58 \pm 0.74	3.22 \pm 0.59	<0.001
2	entropy	3.96 \pm 0.56	3.46 \pm 0.59	3.93 \pm 0.66	3.51 \pm 0.58	<0.001
3	entropy	4.23 \pm 0.53	3.65 \pm 0.59	4.18 \pm 0.65	3.69 \pm 0.59	<0.001
4	entropy	4.39 \pm 0.51	3.80 \pm 0.59	4.35 \pm 0.63	3.85 \pm 0.60	<0.001
5	entropy	4.54 \pm 0.49	3.93 \pm 0.57	4.47 \pm 0.63	3.96 \pm 0.61	<0.001
6	entropy	4.63 \pm 0.48	4.02 \pm 0.59	4.57 \pm 0.62	4.05 \pm 0.60	<0.001
7	entropy	4.72 \pm 0.47	4.08 \pm 0.59	4.64 \pm 0.61	4.12 \pm 0.61	<0.001
8	entropy	4.77 \pm 0.47	4.15 \pm 0.59	4.70 \pm 0.60	4.18 \pm 0.60	<0.001
9	entropy	4.82 \pm 0.46	4.21 \pm 0.59	4.75 \pm 0.59	4.24 \pm 0.60	<0.001
10	entropy	4.87 \pm 0.46	4.26 \pm 0.59	4.79 \pm 0.58	4.28 \pm 0.60	<0.001

Table 3. Post hoc p -values between the phases of the protocol for differences in tone and entropy values with varying lags. P_{12} denotes post hoc p -value between the first baseline phase and heel stimulation phase. P_{13} , post hoc p -value between the first baseline phase and second baseline phase. P_{14} , post hoc p -value between the first baseline phase and heel stick blood drawing phase. P_{23} , post hoc p -value between the heel stimulation phase and second baseline phase. P_{24} , post hoc p -value between the heel stimulation phase and heel stick blood drawing phase. P_{34} , post hoc p -value between the second baseline phase and heel stick blood drawing phase.

	lag	P_{12}	P_{13}	P_{14}	P_{23}	P_{24}	P_{34}
tone	1	<0.001	>0.999	0.015	<0.001	0.052	0.013
	2	<0.001	>0.999	0.001	<0.001	0.008	0.001
	3	<0.001	>0.999	<0.001	<0.001	0.005	<0.001
	4	<0.001	>0.999	<0.001	<0.001	0.003	<0.001
	5	<0.001	>0.999	<0.001	<0.001	0.003	<0.001
	6	<0.001	>0.999	<0.001	<0.001	0.004	<0.001
	7	<0.001	>0.999	<0.001	<0.001	0.005	<0.001
	8	<0.001	>0.999	<0.001	<0.001	0.006	<0.001
	9	<0.001	>0.999	<0.001	<0.001	0.008	<0.001
	10	<0.001	>0.999	<0.001	<0.001	0.008	<0.001
entropy	1	0.002	0.990	0.023	0.005	0.863	0.055
	2	0.001	0.997	0.004	0.003	0.986	0.009
	3	<0.001	0.976	<0.001	<0.001	0.994	0.001
	4	<0.001	0.985	<0.001	<0.001	0.978	0.001
	5	<0.001	0.945	<0.001	<0.001	0.992	0.001
	6	<0.001	0.959	<0.001	<0.001	0.995	<0.001
	7	<0.001	0.931	<0.001	<0.001	0.987	<0.001
	8	<0.001	0.945	<0.001	<0.001	0.996	<0.001
	9	<0.001	0.932	<0.001	<0.001	0.99	<0.001
	10	<0.001	0.939	<0.001	<0.001	0.997	<0.001

(p -values: from lag 1 to lag 10 – <0.001), and Phase 3 and Phase 4 (p -values: lag 1 – 0.013, lag 2 – 0.001, from lag 3 to lag 10 – <0.001). Statistically significant differences were observed also between the two stress phases (Phase 2 and Phase 4, p -values: from lag 2 to lag 10 – <0.01), except at lag 1 ($p = 0.052$).

A similar pattern was observed comparing entropy values among Phases 1 and 2, Phase 1 and Phase 3, as well

as Phase 3 and Phase 2. Phase 3 and 4 did not differ only at lag 1 ($p = 0.055$), while no significant differences were observed at any lag comparing Phases 2 and 4.

The results of ROC analyses are reported as AUC and 95% confidence intervals, between Phases 1 and 2, and Phases 1 and 4 (table 4 and figure 2). Statistically significant differences between the AUC values of tone and entropy, between Phases 1 and 2 were observed at every lag ($p < 0.05$), whereas

Table 4. Average AUC values with 95% confidence interval (CI) obtained using tone and entropy between the first baseline and the stress phases (heel stimulation and heel stick blood drawing). *P*-values were calculated using a two-tailed *t*-test and show the statistical significance of the difference in AUC values obtained using tone and entropy parameters. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 4, heel stick blood drawing phase.

lag	Phase 1 versus Phase 2					Phase 1 versus Phase 4				
	tone		entropy		<i>p</i> -value	tone		entropy		<i>p</i> -value
	AUC	95% CI	AUC	95% CI		AUC	95% CI	AUC	95% CI	
1	0.89	0.80–0.95	0.76	0.63–0.83	<0.000	0.74	0.62–0.82	0.69	0.54–0.80	0.026
2	0.88	0.74–0.95	0.76	0.67–0.87	<0.000	0.74	0.62–0.83	0.72	0.55–0.84	0.127
3	0.89	0.80–0.97	0.78	0.67–0.89	<0.000	0.77	0.68–0.86	0.75	0.62–0.83	0.242
4	0.89	0.70–0.95	0.77	0.63–0.88	0.002	0.78	0.63–0.85	0.76	0.65–0.90	0.450
5	0.89	0.74–0.95	0.76	0.65–0.87	0.002	0.79	0.68–0.89	0.78	0.66–0.90	0.355
6	0.88	0.78–0.97	0.77	0.65–0.88	0.001	0.78	0.66–0.86	0.78	0.68–0.87	0.394
7	0.88	0.72–0.95	0.78	0.60–0.88	0.003	0.80	0.69–0.87	0.78	0.68–0.88	0.393
8	0.87	0.74–0.94	0.77	0.61–0.84	0.021	0.79	0.68–0.88	0.79	0.67–0.86	0.290
9	0.87	0.75–0.95	0.77	0.63–0.88	0.005	0.79	0.64–0.87	0.69	0.54–0.80	0.347
10	0.87	0.72–0.93	0.77	0.58–0.85	0.006	0.79	0.69–0.86	0.72	0.55–0.84	0.375

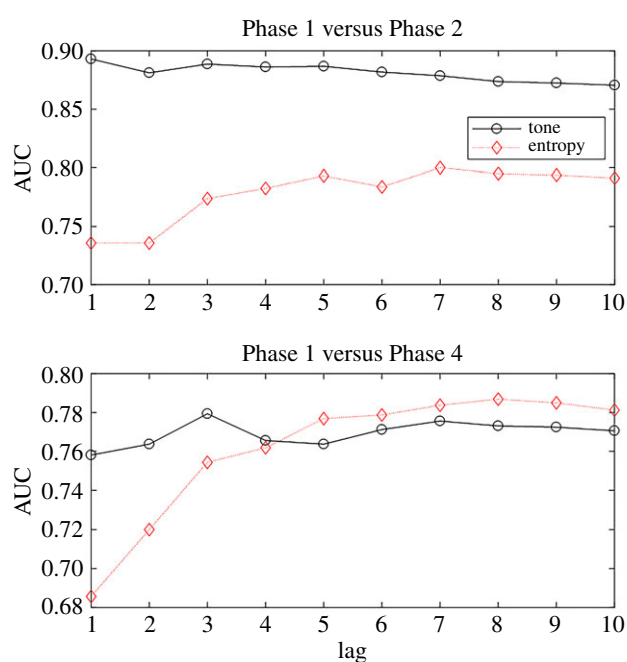


Figure 2. Average AUC value obtained using tone and entropy parameters for Phase 1 versus Phase 2 and Phase 1 versus Phase 4 with varying lag. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 4, heel stick blood drawing phase. (Online version in colour.)

comparing Phase 1 and 4, the AUC values of tone and entropy significantly differed only at lag 1 ($p = 0.026$). The AUC values of tone show consistency comparing Phase 1 and Phase 2, ranging from 0.87 at higher, to 0.89 at lower lags. The obtained AUC values comparing Phase 1 and 4 also showed consistency, ranging from 0.76 at lag 1, 2 and 5, up to 0.78 at lag 7.

In both cases, the discriminating performance of entropy increased with increasing lags, plateauing at lag 5. At higher lag values, entropy showed better performance than tone when comparing Phases 1 and 4. While comparing Phases 1 and 2, AUC values of tone were always greater.

AUC values obtained using both tone and entropy features at each lag using linear regression, showed consistency and high AUC values ($AUC > 0.9$) comparing Phases 1 and 2 (table 5 and figure 3). An increase of AUC with increasing lags was observed comparing Phases 1 and 4 performing best at lags 6 and 9 ($AUC = 0.87$). Significance testing showed better performance of the quadratic model discriminating between Phase 1 and 2, than Phase 1 and 4, at all lags.

4. Discussion

The multi-lag T–E method efficiently shows the changes in the ANS balance that can be used even with short time series. The physiological interpretation of both tone and entropy has been reported in different experimental settings [23,27,32,33]. A lower tone (negative values) indicates the parasympathetic predominance while a higher tone and lower entropy indicate a decrease in the parasympathetic and an increase or sympathetic predominance [23]. Although there is limited research on neonatal pain and stress applying HRV measures, this is the first study investigating multi-lag T–E in a neonatal stress framework [34,35].

It has been previously reported that healthy adults at rest have higher mean levels of entropy and lower and negative mean tone levels at different lags, showing a predominance in the parasympathetic branch of the ANS [22,23]. Our findings are comparable to that of adults, showing the same distribution of mean tone and entropy levels indicating a predominance of parasympathetic activity in term neonates when at rest [36,37].

Compared to the stress phases, at baseline, higher mean entropy values were observed than in the heel stick blood sampling phase, showing a reduction in randomness with increased sympathetic activity. Mean tone values are lower in the baseline phases compared with the stress phases, showing a predominance in parasympathetic activity

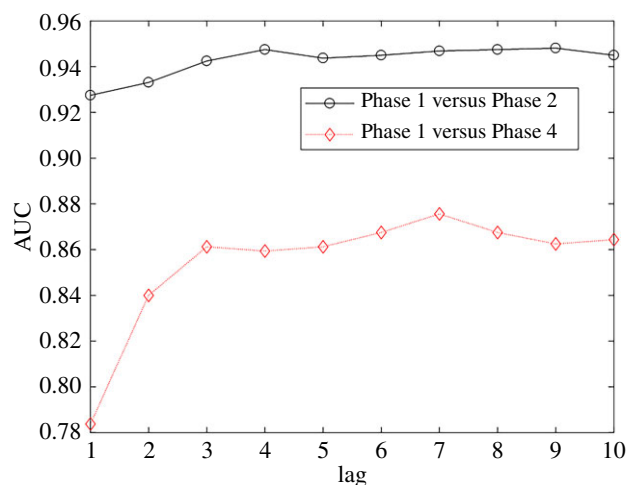


Figure 3. Average AUC value obtained using a linear regression model of both tone and entropy parameters for Phase 1 versus Phase 2 and Phase 1 versus Phase 4 with varying lag. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 4, heel stick blood drawing phase. (Online version in colour.)

Table 5. Average AUC values with 95% confidence interval (CI) obtained using both tone and entropy features at each lag using linear regression (quadratic model). *P*-values were calculated using a two-tailed *t*-test and show the statistical significance of the difference in AUC values obtained for 'Phase 1 versus Phase 2' and 'Phase 1 versus Phase 4' using both tone and entropy parameters. Phase 1, first baseline phase; Phase 2, heel stimulation phase; Phase 4, heel stick blood drawing phase.

lag	Phase 1 versus Phase 2		Phase 1 versus Phase 4		<i>p</i> -value
	AUC	95% CI	AUC	95% CI	
1	0.93	0.85–0.97	0.78	0.62–0.87	<0.001
2	0.93	0.85–0.97	0.84	0.72–0.92	0.001
3	0.94	0.85–0.97	0.86	0.76–0.93	0.001
4	0.95	0.89–0.98	0.86	0.74–0.92	0.001
5	0.94	0.86–0.97	0.86	0.77–0.94	0.001
6	0.95	0.87–0.98	0.87	0.73–0.93	0.001
7	0.95	0.87–0.99	0.88	0.75–0.94	<0.001
8	0.95	0.88–0.98	0.87	0.77–0.93	0.001
9	0.95	0.88–0.98	0.86	0.77–0.93	0.001
10	0.95	0.81–0.98	0.86	0.75–0.93	<0.001

during the baseline phase, and an increase in the sympathetic activity at both stress phases.

Statistically, when considering a time series, high entropy is related to information spread over many states, leading to difficult predictions, due to increased uncertainty or randomness. If we focus on the series of RR intervals, high entropy reflects a healthy responsive ANS. Conversely, a decrease in entropy during the stress phases indicates that the series of RR intervals is less erratic or less random. This is consistent with previous findings, that the HRV of neonates in stress phases results in: (1) an increasing part of mean-reversion effect compared to random fluctuations; (2) an increasing negative scaling exponent, depicting a higher autocorrelation

of the accelerations in the RR_i series. This higher autocorrelation, acting as a fractional integral, smooths the series and thus reduces fluctuations [30,38].

The multi-lag part of our method is intended to take into account long-range influence of the baroreflex. This generalization of the T–E method to lags higher than one is consistent with many other methods that incorporate long-term effects. The particularity of the multi-lag T–E approach compared to fractal methods is that it determines the one lag that is the most significant, independent of any model relating various lags as is often stated in the fractal models. This model-free analysis of the lags therefore seems more general.

Another aim of this study was to estimate the diagnostic performances of tone and entropy and when combining both features. When separately comparing the resting and simulation phase, the performance of tone outperformed entropy, but a slight reversion of performance, especially at higher lags was shown when comparing the resting and blood sampling phase. When combined in a quadratic linear regression model, the AUC values increased in both cases. The lower performance comparing Phase 1 and 4 to Phase 1 and 2 might be related to the duration and type of stress that was applied. When sampling blood, pressure to the heel is routinely applied intermittently; after lancing, pressure to the heel is just enough to collect the required amount of blood to do further laboratory tests and then to stop the bleeding. In Phase 2, intermittent pressure only was applied to the neonates' heel for 90 s while at Phase 4, pressure was applied after lancing for blood collection and for haemostasis. These findings imply that multi-lag T–E may be useful for differentiating the type, duration and intensity of stress in the neonate.

As compared to mature nervous systems, the findings in stressed and resting term neonates match the distributions of tone and entropy to adults without and with cardiac autonomic neuropathy (CAN) [22,23,32]. It is well known that in patients with CAN due to the nerve damage, the parasympathetic activity is diminished and the sympathetic tone predominates [39]. These findings let us conclude that T–E can be used as a surrogate measure of ANS in neonates.

In another recent study, similar to ours though evaluating fetal HRV, the authors indirectly showed the evolution of the ANS [33]. Their results showed that tone increases and entropy decreases at all lags in the late-term fetuses, compared to fetuses with a younger estimated gestational age reflecting the maturation of the sympathetic nervous system as the fetus approaches the delivery period. Our study can be thought as an extension of their work, showing the changes in the ANS in healthy term neonates. The transition from intrauterine to extrauterine life is the most complex adaptation in human life [40]. Significant changes occur in the cardiovascular system during the first hours or perhaps first days of life to a transitional circulation that undergoes further development depending on the different circumstances of extrauterine life. Compared to our study, especially during the baseline phases, at all lags, tone is higher both in the early and late fetuses, while entropy was higher in our study group. Such findings might indicate a maturation both in the sympathetic and parasympathetic branch of the ANS, further showing that in our stress phases, a higher sympathetic response is observed. Although the researchers hypothesized that the maturation of the sympathetic branch might indicate preparation for delivery, transition in the circulation plays a role in ANS balance [40]. The higher dominance of the parasympathetic branch at rest might

imply that at the third day after delivery, the stress response caused during vaginal delivery has diminished.

The observed results of multi-lag T–E should not be limited only to scientific research, but rather they suggest a potential application the real-life NICU environment where vital parameters are routinely measured. Neonates admitted to the NICU are generally suffering from diseases for their specific age group, and may suffer from procedural pain varying from repeated venipunctures, umbilical catheter placements, tracheal suctioning, chest tube placements, lumbar punctures, etc. Owing to the applicability on short-term recordings, baseline values of multi-lag T–E can be easily obtained and applied to real-time measurements. A primer of such application may be useful in the titration of analgesics or sedation medicine, for longer lasting procedures. Similarly, an application of multi-lag T–E can be envisioned in the early onset of diseases with a subtle progression, i.e. necrotizing enterocolitis or perinatal infections, where earlier detection would provide opportunity for intervention or prevention.

References

- Dellinger EH, Boehm FH, Crane MM. 2000 Electronic fetal heart rate monitoring: early neonatal outcomes associated with normal rate, fetal stress, and fetal distress. *J. Obstet. Gynaecol.* **182**, 214–220. (doi:10.1016/S0002-9378(00)70515-1)
- Davis M, Emory E. 1995 Sex differences in neonatal stress reactivity. *Child Dev.* **66**, 14–27. (doi:10.2307/1131187)
- Grunau RE, Holsti L, Haley DW, Oberlander T, Weinberg J, Solimano A, Whitfield MF, Fitzgerald C, Yu W. 2005 Neonatal procedural pain exposure predicts lower cortisol and behavioral reactivity in preterm infants in the NICU. *Pain* **113**, 293–300. (doi:10.1016/j.pain.2004.10.020)
- Sapolsky RM, Romero LM, Munck AU. 2000 How do glucocorticoids influence stress responses? Integrating permissive, suppressive, stimulatory, and preparative actions. *Endocr. Rev.* **21**, 55–89. (doi:10.1210/er.21.1.55)
- Charmandari E, Tsigos C, Chrousos G. 2005 Endocrinology of the stress response. *Annu. Rev. Physiol.* **67**, 259–284. (doi:10.1146/annurev.physiol.67.040403.120816)
- Gunnar MR, Hertzgaard L, Larson M, Rigatuso J. 1991 Cortisol and behavioral responses to repeated stressors in the human newborn. *Dev. Psychobiol.* **24**, 487–505. (doi:10.1002/dev.420240704)
- Wehrwein EA, Orer HS, Barman SM. 2011 Overview of the anatomy, physiology, and pharmacology of the autonomic nervous system. *Compreh. Physiol.* **6**, 1239–1278. (doi:10.1002/cphy.c150037)
- McCorry LK. 2007 Physiology of the autonomic nervous system. *Am. J. Pharm. Educ.* **71**, 78. (doi:10.5688/aj710478)
- Sztajzel J. 2004 Heart rate variability: a noninvasive electrocardiographic method to measure the autonomic nervous system. *Swiss. Med. Wkly.* **134**, 514–522.
- Zipes DP, Wellens HJ. 1998 Sudden cardiac death. *Circulation* **98**, 2334–2351. (doi:10.1161/01.CIR.98.21.2334)
- Bell A, Brown D, Halliday H, McClure G, McReid M. 1989 Meningitis in the newborn—a 14 year review. *Arch. Dis. Child.* **64**, 873–874. (doi:10.1136/ad.64.6.873)
- Gentz J, Persson B, Zetterström O. 1969 On the diagnosis of symptomatic neonatal hypoglycemia. *Acta Paediatrica* **58**, 449–459. (doi:10.1111/j.1651-2227.1969.tb04745.x)
- Thompson AM, Bizzarro MJ. 2008 Necrotizing enterocolitis in newborns. *Drugs* **68**, 1227–1238. (doi:10.2165/00003495-200868090-00004)
- Sabel KG, Wadsworth C. 1979 C-reactive protein (CRP) in early diagnosis of neonatal septicemia. *Acta Paediatrica* **68**, 825–831. (doi:10.1111/j.1651-2227.1979.tb08219.x)
- Kramarić K, Šapina M, Milas V, Milas K, Dorner S, Varžič D, Šerfež J, Adelson PD. 2017 The effect of ambient noise in the NICU on cerebral oxygenation in preterm neonates on high flow oxygen therapy. *Signa Vitae* **13**, 52–56. (doi:10.22514/SV133.062017.11)
- Stone LS, Szyf M. 2013 The emerging field of pain epigenetics. *Pain* **154**, 1–2. (doi:10.1016/j.pain.2012.10.016)
- Carbajal R *et al.* 2008 Epidemiology and treatment of painful procedures in neonates in intensive care units. *JAMA* **300**, 60–70. (doi:10.1001/jama.300.1.60)
- Acharya UR, Joseph KP, Kannathal N, Lim CM, Suri JS. 2006 Heart rate variability: a review. *Med. Biol. Eng. Comput.* **44**, 1031–1051. (doi:10.1007/s11517-006-0119-0)
- Acharya R, Lim C, Joseph P. 2002 Heart rate variability analysis using correlation dimension and detrended fluctuation analysis. *ITBM-RBM* **23**, 333–339. (doi:10.1016/S1297-9562(02)90002-1)
- Camm AJ *et al.* 1996 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* **93**, 1043–1065. (doi:10.1161/01.CIR.93.5.1043)
- Sassi R *et al.* 2015 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* **17**, 1341–1353. (doi:10.1093/europace/euv015)
- Karmakar CK, Khandoker AH, Voss A, Palaniswami M. 2011 Sensitivity of temporal heart rate variability in Poincaré plot to changes in parasympathetic nervous system activity. *Biomed. Eng. Online* **10**, 17. (doi:10.1186/1475-925X-10-17)
- Karmakar CK, Khandoker AH, Jelinek HF, Palaniswami M. 2013 Risk stratification of cardiac autonomic neuropathy based on multi-lag tone–entropy. *Med. Biol. Eng. Comput.* **51**, 537–546. (doi:10.1007/s11517-012-1022-5)
- Claudia L, Oscar I, Héctor PG, Marco VJ. 2003 Poincaré plot indexes of heart rate variability capture dynamic adaptations after haemodialysis in chronic renal failure patients. *Clin. Physiol. Funct. Imaging* **23**, 72–80. (doi:10.1046/j.1475-097X.2003.00466.x)
- Martínez-García P, Lerma C, Infante O. 2012 Baroreflex sensitivity estimation by the sequence method with delayed signals. *Clin. Auton. Res.* **22**, 289–297. (doi:10.1007/s10286-012-0173-7)
- Karmakar C, Jelinek H, Khandoker A, Tulppo M, Makikallio T, Kiviniemi A, Huikuri H, Palaniswami M. 2012 Identifying increased risk of post-infarct people with diabetes using multi-lag tone–entropy analysis. In *Engineering in Medicine and Biology Society (EMBC), 2012 Annual Int. Conf. of the IEEE*, pp. 25–28. Piscataway, NJ: IEEE.
- Oida E, Kannagi T, Moritani T, Yamori Y. 1999 Aging alteration of cardiac vagosympathetic balance

- assessed through the tone–entropy analysis. *J. Gerontol. A Biomed. Sci. Med. Sci.* **54**, M219–M224. (doi:10.1093/gerona/54.5.M219)
28. Léger J, Olivier A, Donaldson M, Torresani T, Krude H, Van Vliet G, Polak M, Butler G. 2014 European Society for Paediatric Endocrinology consensus guidelines on screening, diagnosis, and management of congenital hypothyroidism. *Hormone Res. Paediatrics* **81**, 80–103. (doi:10.1159/000358198)
 29. Tansek MZ *et al.* 2015 Phenylketonuria screening and management in southeastern Europe—survey results from 11 countries. *Orphanet J. Rare Dis.* **10**, 68. (doi:10.1186/s13023-015-0283-0)
 30. Šapina M, Garcin M, Kramarić K, Milas K, Brdarić D, Pirić M. 2017 The Hurst exponent of heart rate variability in neonatal stress, based on a mean-reverting fractional Lévy stable motion.
 31. Shannon CE, Weaver W, Burks AW. 1951 The mathematical theory of communication. *Phil. Rev.* **60**, 398. (doi:10.2307/2181879)
 32. Khandoker AH, Jelinek HF, Moritani T, Palaniswami M. 2010 Association of cardiac autonomic neuropathy with alteration of sympatho-vagal balance through heart rate variability analysis. *Med. Eng. Phys.* **32**, 161–167. (doi:10.1016/j.medengphy.2009.11.005)
 33. Khandoker A, Karmakar C, Kimura Y, Endo M, Oshio S, Palaniswami M. 2015 Tone entropy analysis of foetal heart rate variability. *Entropy* **17**, 1042–1053. (doi:10.3390/e17031042)
 34. Cremillieux C, Makhlouf A, Pichot V, Trombert B, Patural H. 2018 Objective assessment of induced acute pain in neonatology with the Newborn Infant Parasympathetic Evaluation index. *Eur. J. Pain* **22**, 1071–1079. (doi:10.1002/ejp.1191)
 35. Weissman A, Zimmer EZ, Aranovitch M, Blazer S. 2012 Heart rate dynamics during acute pain in newborns. *Pflügers Archiv-Eur. J. Physiol.* **464**, 593–599. (doi:10.1007/s00424-012-1168-x)
 36. Šapina M, Kramarić K, Milas K, Milas V, Vujčić D, Dobrić H, Pirić M, Brdarić D, Pušeljčić S. 2017 Poincaré plot indices as a marker for acute pain response in newborns. *Signa Vitae* **13**, 33–36.
 37. Longin E, Gerstner T, Schaible T, Lenz T, König S. 2006 Maturation of the autonomic nervous system: differences in heart rate variability in premature vs. term infants. *J. Perinat. Med.* **34**, 303–308. (doi:10.1515/JPM.2006.058)
 38. Šapina M, Košmider M, Kramarić K, Garcin M, Pirić M, Milas K, Brdarić D. 2018 Asymmetric detrended fluctuation analysis in neonatal stress. *Physiol. Meas.* **39**, 085006. (doi:10.1088/1361-6579/aad425)
 39. Pop-Busui R. 2010 Cardiac autonomic neuropathy in diabetes: a clinical perspective. *Diabetes Care* **33**, 434–441. (doi:10.2337/dc09-1294)
 40. Hillman NH, Kallapur SG, Jobe AH. 2012 Physiology of transition from intrauterine to extrauterine life. *Clin. Perinatol.* **39**, 769–783. (doi:10.1016/j.clp.2012.09.009)