Heart Rate Variability Among Children With Acquired Brain Injury

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Objective To find evidence of autonomic imbalance and present the heart rate variability (HRV) parameters that reflect the severity of paroxysmal sympathetic hyperactivity (PSH) in children with acquired brain injury (ABI).

Methods Thirteen children with ABI were enrolled and age- and sex-matched children with cerebral palsy were selected as the control group (n=13). The following HRV parameters were calculated: time-domain indices including the mean heart rate, standard deviation of all average R-R intervals (SDNN), root mean square of the successive differences (RMSSD), physical stress index (PSI), approximate entropy (ApEn); successive R-R interval difference (SRD), and frequency domain indices including total power (TP), high frequency (HF), low frequency (LF), normalized HF, normalized LF, and LF/HF ratio.

Results There were significant differences between the ABI and control groups in the mean heart rate, RMSSD, PSI and all indices of the frequency domain analysis. The mean heart rate, PSI, normalized LF, and LF/HF ratio increased in the ABI group. The presence of PSH symptoms in the ABI group demonstrated a statistically significant decline of the SDNN, TP, ln TP.

Conclusion The differences in the HRV parameters and presence of PSH symptoms are noted among ABI children compared to an age- and sex-matched control group with cerebral palsy. Within the ABI group, the presence of PSH symptoms influenced the parameters of HRV such as SDNN, TP and ln TP.

Keywords Brain injuries, Autonomic nervous system, Heart rate, Sympathetic nervous system, Parasympathetic nervous system

INTRODUCTION

The immature brain of children is extremely vulnerable to internal and external insults such as ischemia, hypoxia, traumatic force and infection. Such insults can lead to acquired brain injury (ABI) in children and cause disability or even death. Annually, over 500,000 children between birth through the age of 14 years visit emergency
rooms for ABI across the United States [1]. Among these cases, traumatic brain injury (TBI) is the most common cause of disability and death in children with an estimated prevalence of 475,000 cases per year in the United States [2,3].

Paroxysmal sympathetic hyperactivity (PSH) after ABI is described as an increased body temperature of unknown origin, paroxysmal increase in blood pressure (BP), respiratory rate (RR), and heart rate (HR), and muscle rigidity and hyperhidrosis [4]. Since symptoms of PSH resemble nonspecific symptoms of other general medical conditions such as infection, it is often underestimated and result in unnecessary, repetitive laboratory evaluations. Such abnormal symptoms occur in approximately 8%-10.5% of severe TBI patients during the acute phase [5-7]. Usually, symptoms of PSH subside within the first few weeks after injury; however, a small group of patients continue to experience episodes of PSH [8]. Recently, some studies reported PSH in pediatric patients with hypoxic ischemic encephalopathy [9,10].

Clinicians tend to regard autonomic dysfunction as a minor problem; however, PSH deteriorates the clinical and functional well-being of brain injury patients. Increased morbidity that was caused by abnormal hyperthermia, catabolism and spasticity has been reported in untreated PSH patients [11,12]. Despite increasing clinical reports of this condition, it is still under-recognized; therefore, consensus on the diagnosis, evaluation and treatment of PSH in ABI children has not yet occurred.

In spite of the clinical importance of PSH, it is difficult to evaluate one’s autonomic imbalance. Heart rate variability (HRV) is a non-invasive marker of cardiac autonomic function extracted from continuous electrocardiogram (ECG) recordings, which are also an emerging method to evaluate the sympathetic and parasympathetic balance of the autonomic nervous system (ANS) [13].

The aim of this study was to find evidence of autonomic imbalance and present the HRV parameters that reflect the severity of PSH in children with ABI.

MATERIALS AND METHODS

Subjects
A total of 13 children with ABI below the age of 18 years who had been admitted at the Department of Physical Medicine and Rehabilitation from October 2013 to January 2016 were enrolled in this study. The exclusion criteria were (1) history of cardiac disease, (2) diabetes or other peripheral neurological impairment, (3) patients taking cardiovascular medications, and (4) sepsis or suspicion of significant infection [5,14]. To meet the objectives of this study, 13 age- and sex-matched patients with cerebral palsy were selected as the control group. Patients’ demographics and etiology were obtained from their electrical medical records.

For the purposes of this study, the criteria of PSH that was defined by Baguley et al. [15] was modified. PSH was defined as any simultaneous, paroxysmal increase in at least two out of the four reported clinical data of dysautonomia (i.e., HR, RR, temperature, systolic BP [SBP]), immediately after the onset of dystonia or sweating [15]. HR, RR, SBP, and axillary temperature data were collected at the occurrence of dystonia or sweating three consecutive times in 10-minute intervals and the mean value was calculated. Based on the normal HR and RR values for healthy children from birth to 18 years of age and the abnormal HR and RR criteria of advanced pediatric life support [16], abnormal HR and RR were assessed. Axillary temperature was measured using standardized digital thermometry [17]. According to the nationwide data of normal BP levels of children from 1999 to 2000 from the National Heart, Lung and Blood Institute (NHLBI) in the United States, findings above the 95th percentile are defined as abnormal [18]. Dystonia was defined as sudden occurrence or aggravation of rigidity or decerebrated posturing.

HRV measurement
HRV was analyzed using a SA-6000 device (Medicore Co., Seoul, Korea). HRV recordings were obtained for 5 minutes. Tests were conducted in a noise-free environment after 30 minutes of rest. All study participants attached electric patches to both wrists and the left ankle and remained in a relaxed supine position. In order to avoid the influence of circadian changes in the HRV results, data was collected between 1:00 PM and 3:00 PM in every patient. The room temperature during data collection was 24°C–26°C. ECGs were obtained through 5-minute ECG recordings using personal computers. Abnormal beats, significant pauses, and areas of artifact were automatically rejected by using a computerized algorithm. After filtering, the HRV was automatically analyzed by
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the time and frequency domains. Using the fast Fourier transform (FFT), the HRV results were categorized into different frequency domains [19].

The time domain measures that were analyzed include the following: the mean heart rate, standard deviations of all average R-R intervals (SDNN), square root of the mean squared differences of successive R-R intervals (RMSSD), physical stress index (PSI), approximate entropy (ApEn), and successive R-R interval difference (SRD). The frequency domain measures included the following: total power (TP), low frequency component (LF, 0.04–0.15 Hz), high frequency component (HF, 0.15–0.4 Hz), and low to high frequency ratio (LF/HF ratio). The SDNN reflects every cyclic aspect that was related to the variability during the recording period. High SDNN values indicate irregular variability of the heart rate; while low values indicate a decreased variation of the heart rate despite an environmental change [20]. The RMSSD estimates the HF variations in heart rate, which represents the parasympathetic regulation of the heart [21]. The PSI detects the degree of load and pressure on the ANS control system. The ApEn is a measure of regularity versus randomness and quantifies the unpredictable variation of the heart rate. A low ApEn value indicates a regular signal in the test. The SRD is measured by comparing the initial to the successive part of the total measurement. The full scale value is 1, which reflects the constant status during the measurement. Similar to the SDNN of the time domain, the TP of the frequency domain reflects the variability of the R-R intervals [20]. The LF component of the HRV is mediated by both the sympathetic and parasympathetic nerve activities, but reflects sympathetic activity more dominantly. In contrast, the HF component is mediated almost entirely by the parasympathetic nerve activity. The LF/HF ratio verifies the balance between the sympathetic and parasympathetic nerve activities. The LF and HF values were collected as normalized LF and normalized HF because of time limitations.

Statistical analysis

Data were analyzed with the chi-square test or Fisher exact test for categorical data (i.e., sex, comorbid conditions, criteria of PSH) and the Wilcoxon rank sum test for continuous data (i.e., age, study onset, HRV parameters). The data were analyzed using the Statistical Analysis Software ver. 9.2 (SAS Inc., Cary, NC, USA). All data were reported as a mean (minimum-maximum) for continuous variables and a number (%) for categorical variables. For all analyses, p-values of less than 0.05 were considered statistically significant.

RESULTS

Demographics

This study consisted of 13 children with ABI and 13 sex- and age-matched controls with cerebral palsy. The most common etiology of the ABI group was hypoxic ischemic injury, followed by TBI and neoplasm. Among the control group, 6 were born premature and 7 were born at full-term (Table 1). In the ABI group, the median age was 7.2 years and the median duration from brain injury to HRV measurement was 535 days. The ABI group displayed

Table 1. Etiologies and locations of brain lesions in the ABI and control groups

|                    | ABI group                     | Control group                |
|--------------------|-------------------------------|------------------------------|
| Etiologies         | (n=13)                        | (n=13)                       |
| Hypoxic ischemic   | 7 (53.85)                     | -                            |
| injury             |                               |                              |
| Global injury      | 5                              | -                            |
| Cortical injury    | 1                              | -                            |
| Subcortical injury | 1                              | -                            |
| Traumatic brain    | 4 (30.77)                     | -                            |
| injury             |                               |                              |
| Global injury      | 2                              | -                            |
| Cortical injury    | 1                              | -                            |
| Subcortical injury | 1                              | -                            |
| Neoplasm           | 2 (15.38)                     | -                            |
| Global injury      | 2                              | -                            |
| Cortical injury    | 0                              | -                            |
| Subcortical injury | 0                              | -                            |
| Preterm            | -                              | 6 (46.15)                    |
| Etiologies         | (n=13)                        |                              |
| Periventricular    | -                             | 5                            |
| white matter       |                               |                              |
| lesion             |                               |                              |
| Cortical lesion    | -                             | 0                            |
| Subcortex involvement | -                     | 1                            |
| Term               | -                             | 7 (53.85)                    |
| Etiologies         | (n=13)                        |                              |
| Periventricular    | -                             | 4                            |
| white matter       |                               |                              |
| lesion             |                               |                              |
| Cortical lesion    | -                             | 1                            |
| Subcortex involvement | -                      | 2                            |

Values are presented as number (%). ABI, acquired brain injury.
significantly higher Gross Motor Function Classification System (GMFCS) levels and lower Coma Recovery Scale (CRS) scores than the control group (p<0.0001 and p=0.0001, respectively) (Table 2).

### Table 2. Demographics and comorbid conditions of the ABI and control groups

|                     | ABI group (n=13) | Control group (n=13) | p-value |
|---------------------|------------------|----------------------|---------|
| Age (yr)            | 7.2 (1.8–16.0)   | 7.5 (1.9–16.0)       | 0.8172  |
| Sex                 |                  |                      |         |
| Male                | 8 (61.54)        | 8 (61.54)            |         |
| Female              | 5 (38.46)        | 5 (38.46)            |         |
| Days from brain injury | 535 (49–1,458) | 880 (326–6,177)      | 0.0455* |
| GMFCS               | 5 (4–5)          | 2 (1–3)              | <0.0001*|
| CRS                 | 6 (0–11)         | 23 (23–23)           | <0.0001*|
| History of epilepsy | 9 (69.23)        | 4 (30.77)            | 0.0499* |
| Anticonvulsant      | 10 (76.92)       | 4 (30.77)            | 0.0183* |
| Brain pill          | 2 (15.38)        | 1 (7.69)             | 1.0000  |
| Antispastic medication | 9 (69.23)      | 1 (7.69)             | 0.0013* |
| Complication        |                  |                      |         |
| Cardiac             | 0 (0)            | 1 (7.69)             | 1.0000  |
| Pulmonary           | 1 (7.69)         | 0 (0)                | 1.0000  |
| Endocrine           | 4 (30.77)        | 0 (0)                | 0.0957  |
| Gastrointestinal    | 8 (61.54)        | 0 (0)                | 0.0016* |
| Modified diet       | 3 (23.08)        | 0 (0)                | 0.2200  |
| Genitourinary       | 2 (15.38)        | 0 (0)                | 0.4800  |
| Psychiatric         | 2 (15.38)        | 0 (0)                | 0.4800  |
| Musculoskeletal     | 4 (30.77)        | 4 (30.77)            | 1.0000  |

Values are presented as mean (minimum-maximum) or number (%).

ABI, acquired brain injury; GMFCS, Gross Motor Function Classification System; CRS, Coma Recovery Scale.

*p<0.05 according to the chi-square tests, Fisher exact tests, or Wilcoxon rank sum tests.

The ABI and control group were significantly different in several comorbid conditions: history of epilepsy, usage of anticonvulsant, modified diet and abnormal bladder function. Among the ABI group, 69.23% had a seizure history (p=0.0499), and consequently, 76.92% were on anti-epileptic drugs (p=0.0183). Nine patients (69.23%) were on spasticity medication (p=0.0013) and 8 patients (61.54%) were on a semi-liquid diet via L-tube or gastrostomy tube in the ABI group (p=0.0016). Other comorbid conditions such as complications of the cardiac, pulmonary, endocrine, genitourinary, psychiatric, and musculoskeletal systems demonstrated no significant differences between the two groups (Table 2).

**Comorbid condition**

The ABI and control group were significantly different in several comorbid conditions: history of epilepsy, usage of anticonvulsant, modified diet and abnormal bladder function. Among the ABI group, 69.23% had a seizure history (p=0.0499), and consequently, 76.92% were on anti-epileptic drugs (p=0.0183). Nine patients (69.23%) were on spasticity medication (p=0.0013) and 8 patients (61.54%) were on a semi-liquid diet via L-tube or gastrostomy tube in the ABI group (p=0.0016). Other comorbid conditions such as complications of the cardiac, pulmonary, endocrine, genitourinary, psychiatric, and musculoskeletal systems demonstrated no significant differences between the two groups (Table 2).

**Fig. 1.** Comparison of paroxysmal sympathetic hyperactivity (PSH) symptoms between the ABI and control groups. ABI, acquired brain injury; SBP, systolic blood pressure. *p<0.05 by chi-square test or Fisher exact test.
Comparison of PSH symptoms between the ABI and control groups

PSH was diagnosed in 5 patients (38.46%) in the ABI group; however, no one was diagnosed with such in the control group (p=0.0391). To be specific, five categories were significantly higher in the ABI group: abnormal SBP, abnormal PR, sweating, unexplained fever, and dystonia (p<0.0001, p=0.0016, p=0.0016, p<0.0001, and p=0.0013, respectively) (Fig. 1).

Comparison of HRV between the ABI and control groups

**Time domain analysis**

The statistical analysis demonstrated a statistically significant increase in the mean heart rate of the ABI group compared to the control group (p=0.0119). According to the time domain analysis, the RMSSD and PSI were significantly different between the two groups (p=0.0240 and p=0.0354, respectively) (Table 3).

**Frequency domain analysis**

All indices of the frequency domain analysis demonstrated a statistically significant difference between the two groups. The TP, ln TP, LF, ln LF, HF, ln HF, and normalized HF significantly decreased in the ABI group compared to the control group. Only the normalized LF and LF/HF ratio increased in the ABI group compared to the control group (p=0.0018 and p=0.0018, respectively) (Table 3).

Comparison of HRV between the PSH and non-PSH patients among the ABI group

**Time domain analysis**

Among the ABI groups, 5 patients were diagnosed with PSH (PSH-ABI group), while 8 patients did not meet the criteria of PSH (non-PSH-ABI group). The SDNN values in the PSH-ABI group significantly decreased compared to the non-PSH-ABI group (p=0.0134). The remainder of the time domain analysis resulted in no other significant differences between the two subgroups (Table 4).

**Frequency domain analysis**

The TP and ln TP significantly decreased in the PSH-ABI group compared to the non-PSH-ABI group (p=0.0043, Table 3.

### Table 3. Comparison of HRV between the ABI and control groups

|                          | ABI group (n=13) | Control group (n=13) | p-value |
|--------------------------|------------------|----------------------|---------|
| **Time domain**          |                  |                      |         |
| Heart rate               | 111 (95–149)     | 97 (58–133)          | 0.0119* |
| SDNN (ms)                | 25.916 (9.276–73.36) | 47.447 (21.309–57.680) | 0.1509  |
| RMSSD (ms)               | 17.208 (9.290–48.531) | 42.361 (12.231–61.437) | 0.0240* |
| PSI                      | 129.673 (23.908–606.616) | 37.403 (16.132–204.967) | 0.0354* |
| ApEn                     | 1.032 (0.040–1.309) | 1.104 (0.724–1.249) | 0.9591  |
| SRD                      | 1.077 (0.744–1.418) | 1.000 (0.604–1.222) | 0.1996  |
| **Frequency domain**     |                  |                      |         |
| TP (ms²)                 | 406.574 (56.543–5,423.10) | 1581.000 (314.867–3,167.61) | 0.0312* |
| ln TP                    | 6.008 (4.035–8.598) | 7.366 (5.752–8.061) | 0.0312* |
| LF (ms²)                 | 72.431 (16.917–952.924) | 296.300 (93.167–964.410) | 0.0210* |
| ln LF                    | 4.238 (2.828–6.960) | 5.691 (4.535–6.872) | 0.0210* |
| HF (ms²)                 | 51.063 (1.836–237.086) | 271.360 (94.389–1,062.63) | 0.0009* |
| ln HF                    | 3.933 (0.607–5.468) | 5.603 (4.547–6.969) | 0.0009* |
| Normalized LF            | 69.558 (25.659–90.212) | 48.807 (29.913–64.794) | 0.0018* |
| Normalized HF            | 30.442 (9.788–74.341) | 51.193 (35.206–70.087) | 0.0018* |
| LF/HF ratio              | 2.285 (0.345–9.217) | 0.953 (0.427–1.840) | 0.0018* |

Values are presented as mean (minimum-maximum). HRV, heart rate variability; ABI, acquired brain injury; SDNN, standard deviations of normal to normal R-R intervals; RMSSD, square root of the sum of the square of differences between adjacent R-R intervals; PSI, physical stress index; ApEn, approximate entropy; SRD, successive R-R interval difference; TP, total power; LF, low frequency; HF, high frequency.

*p<0.05 according to the Wilcoxon rank sum test.
Comparison of PSH symptoms between the subgroups based on the location of the brain lesion in the ABI and control groups

The ABI group can be divided into three subgroups according to the location and range of the brain lesion: global (n=9), cortical (n=2), and subcortical injury (n=2) [22]. Similarly, the control group can be divided into three subgroups according to the location of the brain lesion: periventricular (n=9), cortical (n=1), and subcortical lesion (n=3). The occurrence of each PSH symptom and diagnosis of PSH were not significantly different between each of the subgroups based on the location of the brain lesion. A trend of a higher diagnostic rate of PSH was noted among global injury subgroups in the ABI group (Table 5).

DISCUSSION

Childhood ABI results in various forms of disturbances involving motor, sensory, cognitive and behavioral dysfunctions. PSH is also observed after ABI; however, it has
rarely been studied, especially in the pediatric popula-
tion. In spite of the scarcity of clinical reports, recognition 
of PSH is crucial because it is associated with increased 
morbidity rates and poorer outcomes [6,8,23,24].

Many scholars have presented various hypotheses re-
garding the pathophysiology of PSH after brain injury. 
Two mainstream hypotheses have been settled: a theory 
inferring an epileptogenic cause and a theory involv-
ing the disconnection syndrome, which suggests injury to 
an extensive brain lesion from the cerebral cortex, 
diencephalon to the upper brainstem and medulla [5].
Based on the epileptogenic theory, previous studies have 
attempted to detect and treat epilepsy in patients suffer-
ing from PSH, though the results were negative [25-27].
Disconnection syndromes are thought to occur by either 
mechanical damage or a neurotransmitter blockade. One 
study supported the disconnection syndrome theory by 
reporting markedly variable LF/HF ratios and decreased 
LF power in severe TBI patients with PSH compared to 
patients without PSH [5]. In spite of such efforts, direct 
evidence of the disconnection syndrome is still lacking 
[4,28]. The excitatory/inhibitory ratio (EIR) model was 
recently presented as a modified version of the discon-
nection theory to explain diverse severe autonomic over-
activity after brain injury, such as neuroleptic malignant 
syndrome, serotonin syndrome, parkinsonian-hyperpy-
rexia syndrome, malignant hyperthermia, and PSH. In 
this model, the autonomic overactivity is explained by 
the damaged diencephalic/brainstem inhibitory center 
which abnormally converts the non-nociceptive affer-
ent stimuli from the spinal cord to nociceptive input and 
consequently increases the sympathetic excitatory activ-
ity [29]. The aim of this study was to prove the abnormal 
EIR of the ANS and identify the HRV parameters that re-
fect the autonomic dysfunction of ABI children with PSH 
compared to age- and sex-matched control children with 
cerebral palsy.

In this study, the ABI group, which had significantly 
higher GMFCS levels and lower CRS scores, displayed a 
higher rate of abnormal SBP, PR, sweating, unexplained 
fever and dystonia, which resulted in a significantly 
higher diagnostic rate of PSH. Compared to the control 
group, the ABI group revealed significant increases in the 
mean heart rate and PSI, and a significant decrease in TP, 
which emphasizes the reduction of the ANS in a dynamic 
complexity. Moreover, the significant decline of the RMS-SD, LF, HF; normalized HF and significant incline of the 
normalized LF; LF/HF ratio in the ABI group reflected the 
imbalance between the sympathetic and parasympathet-
ic systems. Since the LF reflected the sympathetic activity 
more dominantly, the significantly increased normalized 
LF of the ABI group indicated high sympathetic excitatory 
activity. While HF reflected the parasympathetic activity, 
the significantly decreased HF and normalized HF values 
of the ABI group implied low parasympathetic activity.
The significant increase of the LF/HF ratio of the ABI 
group mirrored the stronger influence of the sympathetic 
power over parasympathetic power. As a result, this study 
demonstrated that the ABI group has higher sympathetic 
excitatory activity and more dominant sympathetic pow-
er than parasympathetic power compared to the control 
group.

Consistent with our results, previous studies involv-
ing adult traumatic brain injury patients demonstrated 
significantly lower LF, TP, and SDNN, and a significantly 
elevated LF/HF ratio [5,30]. One pediatric HRV study of 
boys who suffered severe traumatic brain injury exhib-
ited a significantly higher heart rate at rest and during 
exercise compared to typically-developed controls. This 
study also reported a lower RMSSD of the HRV at rest 
and significantly decreased time-domain values during ex-
ercise only in traumatic brain injury children [31].

For clinical convenience, the measurement of vital 
signs (i.e., HR, RR, SBP, and temperature) that were used 
as the diagnostic criteria of PSH promptly after the onset 
of dystonia or sweating instead of continuous monitoring 
was used. Among the ABI patients, 5 patients (38.46%) 
were diagnosed with PSH. The PSH-ABI group had sig-
nificantly lower values of the SDNN and TP compared to 
the non-PSH-ABI group. Since the SDNN and TP values 
represent the irregular variability of heart rate, decreased 
SDNN and TP indicate a less variable nature of the heart 
rate during environmental change. In short, the ABI 
group has less adaptability of their ANS to a changing 
environment. Although the mean heart rate and LF/HF 
ratio were higher values in the PSH-ABI group, it was not 
statistically significant. The only previous study of HRV 
measurement that compared PSH and non-PSH groups 
in ABI suggested significantly lower LF values in adult 
TBI patients with PSH [5]. This difference may be due to 
the small sample size of the previous and current studies 
(n=16, n=13, respectively). Nonetheless, the overall trend
of the HRV parameters reflected a lack of adaptability of the ANS to a continuously changing environment and a predominance of sympathetic excitatory power over parasympathetic inhibitory power, which supports the EIR model [32].

There are a number of limitations of this study. First, the number of subjects included in this study was relatively small due to the relative scarcity of inpatient, ABI pediatric patients. To overcome this limitation, we conducted a sex- and age-matched case-control study. Second, the sex- and age-specific reference values of the HRV parameters have not been established, especially in children. Since the normal value of the heart rate differs among sex and age in children [33], the normal values of the HRV parameters differ. To produce a more accurate correlation of the HRV parameters and PSH, sex- and age-specific normal values of the HRV parameters are necessary. Third, the results only represented the state of the CNS heart rate control while subjects are at rest. As seen in other dynamic studies conducted at rest and during exercise [28], the HRV parameters change with provocation of the sympathetic tone compared to the resting state, and in patients with PSH, this gap is expected to maximize. Finally, each subject underwent measurement of their HRV at a different period from the onset of disease. As proved by previous studies, patients were more prone to present dysautonomia features during the earlier period after brain injury and exhibited decreased values of the HRV parameters. This different timing of the HRV measurement may influence the overall results of this study. Nevertheless, this study demonstrated that ABI children with PSH presented dominant values of the sympathetic HRV parameters at rest. Further multicenter studies with large sample sizes of HRV measurement at rest and during dysautonomia symptoms in ABI children at the same time period of post-injury is necessary.

In conclusion, the difference in the HRV parameters and presence of PSH symptoms were noted among ABI children compared to an age- and sex-matched control group with cerebral palsy. The presence of PSH symptoms in pediatric ABI patients influenced the parameters of HRV, especially the SDNN and TP values that represent the irregular variability of heart rate.

**CONFLICT OF INTEREST**

No potential conflict of interest relevant to this article was reported.

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