Challenges of deceleration and acceleration capacity of heart rate in patients with acute hemispheric ischemic stroke

Yan-Hong Xu1
Xing-De Wang2
Jia-Jun Yang1
Li Zhou2
Yong-Chao Pan1

1Department of Neurology,
2Department of Cardiology, Shanghai Jiao Tong University Affiliated Sixth People’s Hospital, Shanghai, People’s Republic of China

Background and purpose: Autonomic dysfunction is common after stroke, which is correlated with unfavorable outcome. Phase-rectified signal averaging is a newly developed technique for assessing cardiac autonomic function, by detecting sympathetic and vagal nerve activity separately through calculating acceleration capacity (AC) and deceleration capacity (DC) of heart rate. In this study, we used this technique for the first time to investigate the cardiac autonomic function of patients with acute hemispheric ischemic stroke.

Methods: A 24-hour Holter monitoring was performed in 63 patients with first-ever acute ischemic stroke in hemisphere and sinus rhythm, as well as in 50 controls with high risk of stroke. DC, AC, heart rate variability parameters, standard deviation of all normal-to-normal intervals (SDNN), and square root of the mean of the sum of differences between adjacent normal-to-normal intervals (RMSSD) were calculated. The National Institutes of Health Stroke Scale (NIHSS) was used to assess the severity of stroke. We analyzed the changes of DC, AC, SDNN, and RMSSD and also studied the correlations between these parameters and NIHSS scores.

Results: The R–R (R wave to R wave on electrocardiogram) intervals, DC, AC, and SDNN in the cerebral infarction group were lower than those in controls (P=0.003, P=0.002, P=0.006, and P=0.043), but the difference of RMSSD and the D-value and ratio between absolute value of AC (|AC|) and DC were not statistically significant compared with those in controls. The DC of the infarction group was significantly correlated with |AC|, SDNN, and RMSSD (r=0.857, r=0.619, and r=0.358; P<0.000, P=0.000, and P=0.004). Correlation analysis also showed that DC, |AC|, and SDNN were negatively correlated with NIHSS scores (r=-0.279, r=-0.266, and r=-0.319; P=0.027, P=0.035, and P=0.011).

Conclusion: Both DC and AC of heart rate decreased in patients with hemispheric infarction, reflecting a decrease in both vagal and sympathetic modulation. Both DC and AC were correlated with the severity of stroke.

Keywords: acute ischemic stroke, autonomic dysfunction, deceleration capacity of heart rate, acceleration capacity of heart rate, heart rate variability

Introduction
Stoke has become the leading cause of death in the People’s Republic of China1 and the second most common cause of death worldwide.2 Cardiac autonomic dysfunction is common after stroke, which is correlated with unfavorable outcome.3,4 As an electrophysiological method, heart rate variability (HRV) has been used to assess the autonomic function for decades. Studies have shown that HRV is related to the severity of stroke3 and has some mortality predictive values.6 However, it cannot accurately distinguish the sympathetic and vagal nerve activity. Phase-rectified signal averaging...
(PRSA) is a newly developed technique for detecting cardiac autonomic function, which can quantitatively detect the function of vagal and sympathetic nerves by calculating the deceleration capacity (DC) and acceleration capacity (AC) of the heart, respectively. Its advantages have been shown in the field of cardiology. However, this method has not yet been used in the study of autonomic dysfunction after stroke. In this study, the changes in DC and AC of heart rate and other related parameters were studied in patients with acute cerebral infarction for the purpose of investigating their values in assessing the autonomic function of patients with stroke.

Materials and methods

Subjects

A total of 63 patients (38 men and 25 women; mean age, 71±12 years) with first-ever acute cerebral infarction and sinus rhythm were recruited within 72 hours after stroke. A magnetic resonance imaging (MRI) or computed tomography (CT) scan confirmed the infarction located in the cerebral hemispheres, with 35 in the left hemisphere and 28 in the right. Patients with manifestations of other nervous system lesions and patients with any other diseases or medication known to affect the autonomic nervous system were excluded. Patients with tumor, infection, previous heart, or pulmonary disease were also excluded.

A total of 50 controls (24 men and 26 women; mean age, 68±11 years) were recruited from subjects with high risk of stroke but without history of stroke. Patients who had two or more of the following major risk factors, or one major risk factor, and two or more secondary risk factors were defined as patients with high risk of stroke. Major risk factors include 1) hypertension, 2) hyperlipidemia, 3) diabetes mellitus, 4) age >50 years, and 5) metabolic syndrome; secondary risk factors include 1) atrial fibrillation or heart disease, 2) smoking, 3) family history of stroke or heart attack, 4) obesity, 5) regular alcohol consumption, 6) lack of physical exercise, 7) dietary excess oil, 8) sleep apnea, 9) male, 10) often gums bleeding, teeth loose, or loss, 11) ischemic ophthalmopathy, and 12) sudden deafness. MRI was used in the control group to exclude acute infarction. The other exclusion criteria were the same as for the stroke group.

We obtained oral informed consent from both patients and controls or their clients. This study was approved by the ethics committee of Shanghai Jiao Tong University Affiliated Sixth People’s Hospital. Sex, age, and combined basic diseases and conditions were not statistically different between the two groups (Table 1).

Clinical materials

MRI was used to obtain the side and location information of the infarction. If MRI was contraindicated or unable to be obtained, repeat CT scans were used in order to identify the exact location of infarction. The National Institutes of Health Stroke Scale (NIHSS) was used to record the severity of neurologic deficit. An experienced neurologist was in charge of this work.

Detection of autonomic parameters

All patients underwent a 24-hour Holter monitoring within 1 week after admission. The recordings were analyzed by a DMS Holter system (DMS300-3; Diagnostic Monitoring Software, Stateline, NV, USA). The interferences were excluded, and the events of arrhythmia were tagged artificially when playing back the recordings. The signal-processing technique of PRSA was used to process sequences of R–R intervals obtained from the Holter recordings. This technique provides separate characterizations of deceleration- and acceleration-related modulations, quantified by DC and AC. It is believed that DC reflects vagal activity and AC reflects sympathetic activity of the heart. The computer calculated DC and AC automatically. Meantime, standard deviation of all normal-to-normal intervals (SDNN) and square root of the mean of the sum of the squares of differences between adjacent normal-to-normal intervals (RMSSD) were also calculated through the 24-hour Holter recordings, which

<table>
<thead>
<tr>
<th>Characteristic/parameter</th>
<th>Infarction patients (n=63)</th>
<th>Controls (n=50)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male, n (%)</td>
<td>38 (60.3)</td>
<td>24 (48)</td>
<td>0.191</td>
</tr>
<tr>
<td>Age, years</td>
<td>70.7±11.92</td>
<td>68.2±10.66</td>
<td>0.258</td>
</tr>
<tr>
<td>Hypertension, n (%)</td>
<td>50 (79.4)</td>
<td>35 (70.0)</td>
<td>0.252</td>
</tr>
<tr>
<td>Diabetes mellitus, n (%)</td>
<td>21 (33.3)</td>
<td>12 (24.0)</td>
<td>0.278</td>
</tr>
<tr>
<td>Hyperlipidemia, n (%)</td>
<td>18 (28.6)</td>
<td>16 (32.0)</td>
<td>0.693</td>
</tr>
<tr>
<td>Smoking, n (%)</td>
<td>27 (42.9)</td>
<td>20 (40.0)</td>
<td>0.760</td>
</tr>
<tr>
<td>Regular alcohol consumption, n (%)</td>
<td>4 (6.3)</td>
<td>5 (10.0)</td>
<td>0.506</td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>25.2±4.5</td>
<td>24.7±3.9</td>
<td>0.537</td>
</tr>
<tr>
<td>NIHSS</td>
<td>6.52±5.68</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>R–R intervals, ms</td>
<td>865.3±123.9</td>
<td>931.1±90.8</td>
<td>0.003</td>
</tr>
<tr>
<td>DC, ms</td>
<td>5.55±1.97</td>
<td>6.66±1.63</td>
<td>0.002</td>
</tr>
<tr>
<td>AC, ms</td>
<td>−6.05±2.02</td>
<td>−7.06±1.77</td>
<td>0.006</td>
</tr>
<tr>
<td>SDNN, ms</td>
<td>93.1±31.07</td>
<td>103.92±23.14</td>
<td>0.043</td>
</tr>
<tr>
<td>RMSSD, ms</td>
<td>26.05±12.08</td>
<td>28.24±12.61</td>
<td>0.349</td>
</tr>
<tr>
<td></td>
<td>AC</td>
<td>–DC</td>
<td>0.50±0.07</td>
</tr>
<tr>
<td></td>
<td>AC</td>
<td>/DC</td>
<td>1.159±0.529</td>
</tr>
</tbody>
</table>

Abbreviations: AC, acceleration capacity; DC, deceleration capacity; NIHSS, National Institutes of Health Stroke Scale; RMSSD, square root of the mean of the sum of the squares of differences between adjacent normal-to-normal intervals; SDNN, standard deviation of all normal-to-normal intervals; R–R intervals, R wave to R wave intervals on electrocardiogram; |AC|, absolute value of AC.

Table 1 Comparison between infarction patients and controls
were widely used parameters of HRV and were considered to reflect overall variability and parasympathetic modulation of the heart, respectively. As the values of AC were negative, we used the absolute value of AC (|AC|) in the analyses. We also calculated the D-value between |AC| and DC (|AC|−DC) and the ratio of |AC| and DC (|AC|/DC) to reflect the balance of sympathetic and parasympathetic modulation. The electrophysiological analysts remained blinded to the clinical data during the study period.

Statistical analysis
Mean values (± standard deviation [SD]) were calculated for continuous variables. Distributions of continuous variables were determined by the Shapiro–Wilk test. Group differences were assessed by Student’s t-test or Mann–Whitney U-test. Comparisons of categorical variables were made using chi-square test or Fisher’s exact test. Correlations between different autonomic parameters were calculated using Pearson’s correlation test, and correlations between autonomic parameters and NIHSS scores were calculated using Spearman rank correlation test. Data analysis was performed using SPSS Version 17.0 (SPSS Inc., Chicago, IL, USA). A value of P<0.05 was considered statistically significant.

Results
Comparison of autonomic parameters between infarction patients and controls
Compared with controls, the R–R intervals, DC, AC, and SDNN of the infarction group were lower. But the difference of RMSSD, the D-value between |AC| and DC (|AC|−DC), and the ratio of |AC| and DC (|AC|/DC) were not statistically significant compared with controls (Table 1).

Comparison between left and right hemispheric infarction
Some autonomic parameters of patients with the right hemispheric infarction were slightly lower than that of the left, but the differences were not statistically significant (Table 2).

Correlation analysis
In the infarction group, DC correlated significantly with |AC|, SDNN, and RMSSD. In 63 patients with stroke, NIHSS scores ranged from 0 to 21 (6.52±5.68). There is a negative correlation between NIHSS and DC, |AC|, and SDNN. No significant correlation was identified between NIHSS scores and RMSSD, the D-value between |AC| and DC, and the ratio of |AC| and DC (Table 3; Figures 1–3).

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Left infarction (n=35)</th>
<th>Right infarction (n=28)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>NIHSS</td>
<td>7.60±6.44</td>
<td>5.18±4.30</td>
<td>0.080</td>
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<tr>
<td>R–R intervals, ms</td>
<td>869.4±119.2</td>
<td>860.2±150.5</td>
<td>0.789</td>
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<tr>
<td>DC, ms</td>
<td>5.66±1.81</td>
<td>5.42±1.29</td>
<td>0.627</td>
</tr>
<tr>
<td>AC, ms</td>
<td>–6.30±2.00</td>
<td>–5.75±2.05</td>
<td>0.288</td>
</tr>
<tr>
<td>SDNN, ms</td>
<td>93.40±29.98</td>
<td>92.82±32.94</td>
<td>0.942</td>
</tr>
<tr>
<td>RMSSD, ms</td>
<td>27.49±12.91</td>
<td>24.25±10.93</td>
<td>0.295</td>
</tr>
<tr>
<td></td>
<td>AC</td>
<td>−DC</td>
<td>0.63±0.962</td>
</tr>
<tr>
<td></td>
<td>AC</td>
<td>/DC</td>
<td>1.13±2.043</td>
</tr>
</tbody>
</table>

Abbreviations: AC, acceleration capacity; DC, deceleration capacity; NIHSS, National Institutes of Health Stroke Scale; RMSSD, square root of the mean of the sum of the squares of differences between adjacent normal-to-normal intervals; SDNN, standard deviation of all normal-to-normal intervals; R–R intervals, R wave to R wave intervals on electrocardiogram; |AC|, absolute value of AC.

Discussion
Patients with stroke often complicate with cardiac autonomic dysfunction, which can lead to a variety of cardiac arrhythmias, T wave change, myocardial infarction, and even sudden death. Cardiac involvement is an important cause of death after stroke. Sympathetic hyperactivity and decrease in parasympathetic activity caused by stroke may be the reason of arrhythmia and sudden death. Studies found that cardiac autonomic dysfunction was related to the location and severity of stroke and may be related to the unfavorable outcome of stroke.

HRV has been used to detect the cardiac autonomic modulation for decades. However, it cannot distinguish the sympathetic and vagus nerve activity. In addition, there is no HRV parameter directly reflecting sympathetic modulation. Plasma catecholamine concentrations are usually used to measure the function of sympathetic nerve. However, plasma catecholamine concentrations are affected by many factors, such as the release, distribution, metabolism, and excretion of amines, and they are also affected by circadian rhythm.
PRSA is a new technique for detecting autonomic function. It can detect the AC and DC of the heart directly through analyzing the overall trend of 24-hour heart rate, and thus it can quantitatively detect the sympathetic activity and vagus activity, respectively, at the same time. This technique has been used mainly in the field of cardiovascular medicine, but not yet in the analysis of autonomic dysfunction after stroke.

In our study, we found that DC and AC as well as SDNN were all lower than controls, reflecting both sympathetic and vagal modulation loss in patients with hemispheric infarction. DC and AC were correlated with each other strongly, and the D-value and the ratio of absolute value of AC and DC had no significant difference compared with controls, reflecting that there was no major change in sympathovagal balance in patients with stroke. The loss of parasympathetic modulation is consistent with HRV studies. However, the decrease in AC reflects a decrease in sympathetic activity, which is inconsistent with studies concluding that the sympathetic activity of stroke increased through the detection of catecholamine concentrations. In the clinic, increased blood pressure, heart rate, and other characteristics of enhanced sympathetic activity are common in patients with cerebral infarction. In our study, we also found that R–R intervals of patients with infarction were shorter than those in controls, which might be a reflection of sympathetic hyperactivity.

Our findings demonstrate a phenomenon that both the vagal and sympathetic modulation decrease after stroke, but the combined effects of the two result in predominant sympathetic activity. This phenomenon could be explained by the fact that the effect of heart rate increasing by decreased vagal activity was larger than the effect of heart rate decreasing by decreased sympathetic activity. The sympathetic activity is relatively predominant rather than absolutely predominant.

The study results were not conclusive of whether the lateralization and location of infarction were associated with autonomic derangement. Most studies found that if the infarctions are located in the right hemisphere especially with insula involvement, the cardiac autonomic dysfunction was more pronounced. In this study, autonomic parameters of patients with left and right cerebral infarction were compared. Although the autonomic parameters of patients with right hemispheric infarction were slightly lower than those of the left hemisphere, the differences were not statistically significant. Due to limited sample size, patients with the insula involvement were not analyzed separately. A further study of a larger number of cases is needed to search for the effect of lateralization and location of infarction on the autonomic nervous system.

In this study, the NIHSS score was used to assess the severity of stroke. Correlation analysis showed that DC, absolute value of AC, and SDNN were negatively correlated with NIHSS scores, which indicates a higher risk of autonomic complications in patients with more severe stroke. Although significant correlations between autonomic parameters and NIHSS scores were found, the \( r \) values were rather low. We assume that autonomic function is affected by many factors.
such as age, sex, and combined diseases, which do not influence the NIHSS scores directly. The infarction location and lateralization also have different effects on autonomic modulation and NIHSS scores.

Declining autonomic modulation predicts poor outcomes in many diseases, such as myocardial infarction,20 chronic heart failure,21 and ischemic stroke.2 Studies also showed that vagus nerve had protective effects.22,23 The decline of vagal modulation may decrease this protective effect. DC, which is a direct reflection of vagal modulation, has shown its value in predicting mortality after myocardial infarction.2 Our study has also shown a decline in DC in patients with hemispheric infarction. Further studies for the value of DC in prognosis prediction in patients with stroke are deserved.

Conclusion
Both DC and AC decrease in patients with acute hemispheric infarction, reflecting the loss of both parasympathetic and sympathetic modulation after stroke. The clinical manifestations of hyperactivity of sympathetic nerve after stroke are possibly a reflection of the relative increase of sympathetic activity caused by more decline of vagal activity. Both DC and AC were correlated negatively with the severity of stroke. For the protective effect of vagal nerve, a decline of vagal modulation might have some predictive values for unfavorable outcomes after stroke. Further studies for the predictive value of DC and AC for the prognosis of stroke are deserved.

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Disclosure
The authors report no conflicts of interest in this work.

References