Research Article

Differential Temporal Evolution Patterns in Brain Temperature in Different Ischemic Tissues in a Monkey Model of Middle Cerebral Artery Occlusion

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Brain temperature is elevated in acute ischemic stroke, especially in the ischemic penumbra (IP). We attempted to investigate the dynamic evolution of brain temperature in different ischemic regions in a monkey model of middle cerebral artery occlusion. The brain temperature of different ischemic regions was measured with proton magnetic resonance spectroscopy (1H MRS), and the evolution processes of brain temperature were compared among different ischemic regions. We found that the normal (baseline) brain temperature of the monkey brain was 37.16°C. In the artery occlusion stage, the mean brain temperature of ischemic tissue was 1.16°C higher than the baseline; however, this increase was region dependent, with 1.72°C in the IP, 1.08°C in the infarct core, and 0.62°C in the oligemic region. After recanalization, the brain temperature of the infarct core showed a pattern of an initial decrease accompanied by a subsequent increase. However, the brain temperature of the IP and oligemic region showed a monotonously and slowly decreased pattern. Our study suggests that in vivo measurement of brain temperature could help to identify whether ischemic tissue survives.

1. Introduction

An elevation in brain temperature is common in acute ischemic stroke and is associated with a worse outcome [1–3]. One potential explanation is that pyrexia increases the brain temperature and the brain metabolic rate and could result in a more rapid exhaustion of limited energy and oxygen supplies and the increased production of free radicals and other toxic substances in ischemic tissue [4, 5]. For different blood flow and metabolic states, brain temperature could be different in different regions of ischemic tissue, that is, the infarct core, ischemic penumbra (IP), and the oligemic region.

Conventional methods of measuring brain temperature by invasive probes may monitor only one or a few locations simultaneously and cursorily, and they are difficult to be accepted clinically [6, 7]. Several magnetic resonance parameters can be used for the noninvasive measurement of regional temperatures, including the diffusion coefficient, the longitudinal relaxation time constant (T1), and the proton resonance frequency (PRF), and the latter is more popular [8, 9]. Magnetic resonance (MR) spectroscopic techniques are, in principle, capable of estimating absolute temperatures. Therefore, it is possible to more accurately measure brain temperature in different regions of ischemic lesions, especially in the IP.

Many experimental and clinical studies have focused on brain temperature in the fields of cerebral infarction measured by magnetic resonance spectroscopy (MRS) [6, 10, 11]. The brain temperature is elevated in the ischemic brain
sooner after stroke, prior to any increase in body temperature, and is significantly higher in probable penumbral tissue than in the infarct core or normal brain [2, 12]. A previous study [13] found that there was a significant difference in brain temperature between ischemic tissue and the contralateral normal hemisphere in patients with middle cerebral artery occlusion. Additionally, the brain temperature of ischemic tissue could increase with the enlargement of lesion volume. However, there have been no reports on the detailed evolution of brain temperature in different regions of ischemic lesion at the early stage of onset.

In this work, we modified equations to measure brain temperature based on MRS PRF techniques in a physiological solution and used the final equation to calculate the brain temperature of different regions in ischemic lesion before and after reperfusion, especially the IP, in monkey middle cerebral artery occlusion (MCAO) models.

2. Materials and Methods

2.1. Animal Model. The experiment was approved by the Animal Care and Use Committee of Tianjin Medical University. Six mature male monkeys (Cynomolgus macaque) with a mean weight of 8.6 kg (range from 8 to 10 kg) were supplied by Experimental Animal Center of Beijing Military Medical Academy.

All animals were fasted for 12–15 hours before the experiment. Anesthesia was induced by ketamine (0.25 mg intramuscular) and atropine (8–10 mg/kg intramuscular) followed by 100 mg ketamine and 50 mg diazepam in a 500 mL NS intravenous drip. Physiological parameters, such as respiration, heart rate, body temperature, and blood pressure, were monitored.

The MCAO model was established through an autotthrombus interventional method [14]. A one-inch incision was made in the right femoral groove, and the femoral artery and vein were isolated. The right femoral artery was punctured, and a 5 French catheter sheath was inserted. The 5 French catheter was introduced over a guide wire and navigated in a common carotid artery under fluoroscopy in a common carotid artery. A 1.7 French microcatheter (Prowler 10 with two markers, Cordis Corporation, 0.55 mm outside diameter) was introduced over a transcend microwire (Cordis). The microcatheter and the microguide wire were navigated into a selected branch of the middle cerebral artery (MCA) and advanced into a wedged position. The animals were maintained on a slow, continuous, heparinized drip infusion (100 U/hour) during the ischemia period. After one hour of ischemia, the microguide wire was removed, and the animal was analyzed by a magnetic resonance imaging (MRI) scanner. If cerebral ischemia was confirmed by diffusion-weighted imaging (DWI), the microcatheter was removed, and 250,000 units of urokinase in 50 mL normal saline (NS) was infused with a high-pressure injector (2 mL/min). Effective reperfusion was confirmed by angiograms.

2.2. MRI and MRS Examination. MRI and MRS were performed with a GE 1.5T Twin Speed Infinity with Excite I magnetic resonance system 1 h after occlusion and 1 h, 3 h, 6 h, 12 h, and 24 h after recanalization. In addition, MRS was performed before occlusion to acquire normal brain temperature. We performed DWI, perfusion-weighted imaging (PWI), and T<sub>2</sub>-weighted imaging (T<sub>2</sub>WI) with a head coil. The T<sub>2</sub>WI parameters were the following: repetition time (TR) = 4000 ms; echo time (TE) = 106.4 ms; averages = 2; slice thickness = 2.5 mm; gap = 0 mm; field of view (FOV) = 18 cm × 18 cm; and matrix = 288 × 256. DWI was obtained using a single-shot spin-echo echo planar sequence with TR = 6000 ms; TE = 96.8 ms; averages = 2; b = 1000 s/mm<sup>2</sup>; FOV = 18 cm × 18 cm; and matrix = 128 × 128. PWI was obtained using a single-shot gradient-recalled echo planar imaging T<sub>2</sub>WI sequence, which is, dynamic susceptibility contrast (DSC) MRI, with TR = 2000 ms; TE = 80 ms; average = 1; FOV = 18 cm × 18 cm; and matrix = 128 × 128. A Gd-DTPA bolus (0.1 mmol/kg) was administered with a power injector at a rate of 2 mL/s, and 10 mL of isotonic saline was injected to wash the pipe. A total of 330 slices were acquired.

We used multivoxel point-resolved-spectroscopy- (PRESS-) localized proton MRS with the voxel grid centered on the slice showing the maximum ischemic lesion on DWI. The imaging parameters were the following: TR = 1500 ms, TE = 135 ms, FOV = 18 cm × 18 cm, matrix = 24 × 24, slice thickness = 10 mm, and total acquired time = 9′60′′. The voxel grid (1 mL) was carefully placed within the brain to include as much of the visible ischemic lesion ipsilateral and contralateral normal brain as possible and to avoid contamination of the spectra with the lipid signal from bone marrow or subcutaneous fat. Almost thirty available voxel grids were obtained. We collected standard three-pulse chemical shift-selective (CHESS) water suppression and nonsuppression imaging. The spectroscopic data of water suppression were obtained with a 99% suppression level.

2.3. Brain Temperature Calculation. Spectroscopic data were analyzed with the Java-based magnetic resonance user interface (jMRUI, http://www.mrui.uab.es/mrui/) package, spectroscopic time domain analysis software. Initially, the zero-order phase correction of the water proton peak (effectively bringing water to a chemical shift of 4.70 ppm) was performed, and the residual water signal was then removed using the Hanckel-Lanczos Singular Value Decomposition (HLSVD) method. Spectroscopic data were Fourier transformed for display and visual quality control purposes using the Advanced Method for Accurate Robust and Efficient Spectral Fitting (AMARES) algorithm within the jMRUI package. Gaussian components were modeled in the time domain, including choline (Cho), creatine (Cr), N-acetyl aspartate-containing compounds (NAA), and lactate (Lac). The chemical shifts (i.e., frequency) of the fitted metabolite peaks were reported to a precision of 0.001 ppm to confirm the chemical shifts of NAA. Spectra were automatically discarded if the fitted line widths were less than 1 Hz or greater than 10 Hz. We inspected the spectroscopic data visually and discarded voxels laying on the edges of the PRESS excitation region, voxels containing cerebrospinal
fluid (CSF) and those with poor quality, for example, having a badly elevated baseline or containing spurious peaks.

Temperature $T$ was derived from the chemical shift of water ($CS_{H2O}$) using the following equation:

$$T = T_{ref} + k(CS_{H2O} - CS_{ref}).$$

(1)

$CS_{ref}$ is the (temperature-independent) chemical shift of a reference compound, $k$ is the coefficient of proportionality, and $T_{ref}$ is the reference temperature. Temperature-dependent changes in hydrogen bonding cause the water chemical shift to vary linearly with temperature at 0.01 ppm per °C. In our scanner, with 4.7 ppm as the chemical shift of water and 37°C as the reference temperature, (1) was changed to the following:

$$T = 37 + 100(CS_{NAA} - CS_{NAAref}).$$

(2)

$CS_{NAA}$ is the apparent chemical shift of NAA, and $CS_{NAAref}$ is the reference chemical shift of NAA. We measured the temperature of a physiological solution (NAA 12.5 mMol, Cr 10.0 mMol and Cho 3.0 mMol) with constant temperature equipment to acquire the $CS_{NAAref}$, which could be fit to the MR scan. There was a linear correlation between the PRF and the temperature of the solution ranging from 34°C to 44°C, and the value of $CS_{NAAref}$ was 2.039 ppm by this method. Therefore, the following was used to calculate the brain temperature of each voxel grid in our study:

$$T = 37 + 100(CS_{NAA} - 2.039).$$

(3)

### Table 1: Normal brain temperature of monkeys.

| No. | Left hemisphere | Right hemisphere |
|-----|----------------|-----------------|
| 1   | 37.05 (0.26)   | 37.07 (0.27)    |
| 2   | 37.13 (0.36)   | 37.11 (0.38)    |
| 3   | 37.26 (0.26)   | 37.28 (0.23)    |
| 4   | 37.12 (0.24)   | 37.15 (0.25)    |
| 5   | 37.23 (0.18)   | 37.24 (0.21)    |
| 6   | 37.16 (0.28)   | 37.17 (0.24)    |
| Total | 37.16 (0.21) | 37.17 (0.23) |

Note. Data are expressed by means (standard deviation).

### 3. Results

#### 3.1. Normal Brain Temperature of the Monkey.

The normal (baseline) brain temperature of the six monkeys was measured before the operation (Table 1). The mean baseline brain temperature was 37.16°C in left hemispheres, 37.17°C in right hemispheres, and 37.16°C in bilateral hemispheres. There was no significant difference in the baseline brain temperature between the bilateral hemispheres by paired t-test ($t = -1.659, P > 0.05$).

#### 3.2. Evolution of Brain Temperature in Monkey MCAO and Reperfusion Models.

MCAO models were successfully established in four out of the six monkeys, including one right and three left sides. Two monkeys were dead at 2 and 14 hours after reperfusion, respectively, probably because of large area of hemorrhagic infarction. And their data were not included in this study. Table 2 shows the brain temperature of different ischemic tissues and the contralateral hemisphere at the artery occlusion stage (0 h) and different time points of the reperfusion stage. At the artery occlusion stage, the mean brain temperature of ischemic tissue was 1.16 ± 0.15°C higher than the baseline brain temperature. The increase in brain temperature was region dependent, with 1.72 ± 0.08°C in the IP, 1.08 ± 0.19°C in the infarct core, and 0.62 ± 0.14°C in the oligemic region compared to the basal brain temperature. In the reperfusion stage, the evolution of brain temperature differed in the infarct core, IP and oligemic region (Figure 2). After recanalization, the brain temperature of the infarct core showed a pattern of an initial decrease accompanied by a subsequent increase. However, the brain temperature of the IP and oligemic region showed a monotonously and slowly decreased pattern. The brain temperature of the contralateral hemisphere increased slightly (0.23°C) compared to the baseline brain temperature.

Table 3 shows the differences in brain temperature between different ischemic tissues and the contralateral hemisphere. There were significant differences in brain temperature between the infarct core and contralateral hemisphere at 0 h, 1 h, 12 h, and 24 h, between the IP and contralateral hemisphere within 6 h, and between the oligemic region and the contralateral hemisphere within 3 h ($P < 0.05$), while no significant differences were found between the infarct core and contralateral hemisphere at 3 h and 6 h.
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Figure 1: Definition of different ischemic regions. DWI (a) and MTT (c) at the artery occlusion stage and T2WI (b) at 24 h are used to define the infarct core, IP, and oligemic region. On the localized DWI of MRS (d), overlapped imaging shows the infarcted core (white), IP (yellow), and oligemia (red).

Table 2: The brain temperature of different ischemic tissues and contralateral hemisphere at artery occlusion stage (0 h) and different time points of reperfusion stage in monkey MCAO model.

| Time after modeling (hours) | Infarct core | IP | Oligemia region | Contralateral hemisphere |
|-----------------------------|--------------|----|-----------------|--------------------------|
| 0                           | 38.26 (0.27) | 38.90 (0.19) | 37.80 (0.14) | 37.46 (0.14) |
| 1                           | 37.81 (0.26) | 38.34 (0.38) | 37.92 (0.34) | 37.37 (0.46) |
| 3                           | 37.21 (0.36) | 38.29 (0.23) | 37.79 (0.33) | 37.41 (0.37) |
| 6                           | 37.26 (0.29) | 38.22 (0.25) | 37.59 (0.35) | 37.31 (0.32) |
| 12                          | 38.48 (0.17) | 37.56 (0.28) | 37.49 (0.38) | 37.33 (0.39) |
| 24                          | 38.63 (0.19) | 37.47 (0.35) | 37.29 (0.35) | 37.28 (0.45) |

Note. Data are expressed by means (standard deviation).

between the IP and contralateral hemisphere at 12 h and 24 h, or between the oligemic region and contralateral hemisphere at and after 6 h ($P > 0.05$).

3.3. Differences in Brain Temperature among Different Ischemic Regions. One-way ANOVA was performed on the brain temperature of different ischemic tissues at different time points (Table 4). There were significant differences in the brain temperature among different ischemic tissues at each time point ($F$ values were 27.64, 8.32, 26.46, 21.55, 27.64, and 47.90 at 0 h, 1 h, 3 h, 6 h, 12 h, and 24 h, resp., $P < 0.05$). Bonferroni’s method was used for post hoc comparisons.
between the infarct core and oligemic region at 1 h and 6 h between the IP and oligemic region at 12 h and 24 h or infarct core and oligemic region at 0 h, 3 h, 12 h, and 24 h the IP and oligemic region within 6 h, and between the infarct core and IP at each time point, between the infarct core and oligemic region at 0 h, 3 h, 12 h, and 24 h (especially CS NAAref) are dependent on the MR scanner and sequence parameters, the temperature equation can be modified based on in vivo models or healthy volunteers [18]. In our study, we used a physiological solution within a constant temperature to acquire the CS NAAref. Therefore, the CS NAAref was more accurate to calculate the brain temperature.

There are also several limitations regarding MRS thermometry. The signal intensity from NAA as a chemical shift reference should be high enough for calculating the temperature. However, NAA decreases gradually after a stroke; as a consequence, studies of the temperature of ischemic brain tissue by MRS should be limited in acute stroke [18]. Another limitation is the environmental temperature of the MR scanner. In most scanners, the temperature should be 18–20 °C perennially. The brain temperature would be lower after a long-term examination. A previous study [12] found that there was a statistically significant reduction in temperature of 0.09°C per scan (P = 0.0001) across four sequential MRS scans on 4 normal volunteers, presumably due to cooling by the air current in the bore of the magnet (or relative cerebral inactivity during the scan). In our study, the MR examination was arranged as DWI, T2WI, PWI, and MRS for 15 min for all of the time points, which can at least partly reduce the effect of environmental temperature on the longitudinal changes of brain temperature.

There were significant differences in brain temperature between the infarct core and IP at each time point, between the IP and oligemic region within 6 h, and between the infarct core and oligemic region at 0 h, 3 h, 12 h, and 24 h (P < 0.05), while there were no significant differences between the IP and oligemic region at 12 h and 24 h or between the infarct core and oligemic region at 1 h and 6 h (P > 0.05).

4. Discussion

Previous studies of brain temperature were limited by the reliance on invasive measurement techniques, such as probes inserted in brain tissue [15–17]. In this situation, the brain temperature could be lower than the actual temperature if the brain surface was exposed to cooler environmental air. Invasive techniques may inevitably induce local microlesions and inflammatory responses adjacent to probes, which might affect brain temperature. Infrared thermometry is a noninvasive method, but it fails to measure the deep brain temperature due to the limited depth penetration and cannot make an accurate localization because of the limited spatial resolution of this method. MRS is a validated method for noninvasively measuring brain temperature using the principle that the water frequency shift relative to N-acetyl aspartate is temperature dependent [8, 10, 18]. Moreover, MRS thermometry could avoid most of the above-mentioned weaknesses of other methods.

Because equations for calculating brain temperature (especially CS NAAref) are dependent on the MR scanner and sequence parameters, the temperature equation can be modified based on in vivo models or healthy volunteers [18]. In our study, we used a physiological solution within a constant temperature to acquire the CS NAAref. Therefore, the CS NAAref was more accurate to calculate the brain temperature.

The perfusion state is critical for the temperature of the brain tissue; for example, reduced blood flow could impair heat exchange and result in a higher temperature in ischemic tissue [2]. According to the perfusion state theory, the brain temperature in the infarct core should increase the most because it has the lowest blood flow. However, we found that the highest increase in brain temperature was in the IP, followed by the infarct core and the oligemic region at the artery occlusion stage in the monkey MCAO model. This finding suggests that the perfusion state was not the only factor for brain temperature elevation in ischemic tissue, and other factors might play a role.

The brain tissue temperature is mainly determined by two processes, heat production and heat radiation. In the infarct core, reduced blood flow induces the reduction of heat radiation and increased anaerobic metabolism induces the increase in heat production, which may both result in the
Table 4: ANOVA among brain temperature of different ischemic tissues and contralateral hemisphere in monkey MCAO model (P value).

| Time (h) | Infarct core and IP | IP and oligemia region | Infarct core and oligemia region |
|---------|---------------------|------------------------|---------------------------------|
| 0       | 0.032*              | 0.007*                 | 0.005*                          |
| 1       | 0.004*              | 0.005*                 | 0.287                           |
| 3       | 0.003*              | 0.013*                 | 0.011*                          |
| 6       | 0.001*              | 0.011*                 | 0.346                           |
| 12      | 0.001*              | 0.078                  | 0.035*                          |
| 24      | 0.001*              | 0.135                  | 0.024*                          |

Note. *represents $P < 0.05$.

Figure 3: Evolution of rCBF and rMTT in different ischemic regions at the artery occlusion stage (0 h) and reperfusion stage in the monkey MCAO model.

elevated brain temperature. In IP tissue, reduced blood flow may induce the reduction of heat radiation, while an increase in heat production is induced by the subsequent events, including both aerobic and anaerobic metabolism [19–22], inflammation and the release of excitatory amino acids [23, 24]. These complex processes may account for the higher elevated brain temperature in the IP than in the infarct core or oligemic regions at the artery occlusion stage. In addition, uncoupling protein 2 (UCP-2) is a natural neuroprotective factor in the human ischemic brain [25–28]. UCP-2 upregulation may regulate ATP synthesis by uncoupling oxidation from phosphorylation, thus dissipating energy as local heat, and may simultaneously be responsible for the early rise in lesion brain temperature after stroke.

After recanalization, the brain temperature of the ischemic tissue was decreased with time, which can be mainly explained by the changes of the perfusion states in these brain regions (Figure 3). In the infarct core, a temporary and rapid increase in rCBF and shortening of the rMTT, that is, overperfusion, result in the clearance of accumulated heat, which can account for the sharply decreased brain temperature. However, after 6 h, both decreased rCBF and delayed MTT lead to the reincrease in the temperature. In particular, at 12 h after recanalization, the brain temperature of the infarct core was even higher than the IP and oligemic region and was accompanied by the lowest rCBF and highest rMTT in the infarct core. However, the brain temperature of the IP and oligemic region recovered to normal levels as in the contralateral hemisphere without repeated elevation, suggesting that the cells in these regions were viable or salvaged.

The brain temperature in the contralateral hemisphere (which may be more closely related to body temperature) is slightly elevated after a stroke, reflecting increased neuronal activity in response to the ischemia [2]. Therefore, in our study, the brain temperature in the ischemic region was evaluated by comparison with the baseline brain temperature, which is more reasonable than the previously used method that regarded the brain temperature of the contralateral hemisphere as a control state.

In the present study, we focused on detailed changes in the brain temperature of different ischemic tissues before and after reperfusion by $^1$H MRS. We found that the IP and infarct core showed different evolution patterns of brain temperature, which may provide us with a method to discriminate the IP from the infarct core. However, our study also had some limitations. The first limitation is that
the classification of different ischemic tissues in our study was based on DWI and PWI at the artery occlusion stage and T$_2$WI at 24 h of the reperfusion stage. IP was defined as low-perfusion tissue with high signal intensity on DWI but normal intensity on T$_2$WI, but this definition may miss part of the IP tissue peripheral to the abnormal DWI regions [29–31]. The second limitation is the relatively small sample size of our study. The last limitation is that we do not know whether the results obtained from the monkey MCAO model can be generalized to human stroke patients. Further studies are needed to overcome the above limitations and to clarify the mechanism underlying the evolution of the brain temperature of ischemic tissue.

5. Conclusion

After ischemic stroke, the infarct core and IP have different evolution patterns of brain temperature, which suggests that in vivo measurement of brain temperature could help to identify whether ischemic tissue survives.

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