Research Article

Blood-Oxygenation-Level-Dependent-(BOLD-) Based R2′ MRI Study in Monkey Model of Reversible Middle Cerebral Artery Occlusion

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Objective. To investigate the value of BOLD-based reversible transverse relaxation rate (R2′) MRI in detecting ischemic penumbra (IP) in a monkey model of reversible middle cerebral artery occlusion (MCAO) and time evolution of relative R2′ in infarcted core, IP, and oligemia.

Materials and Methods. 6 monkeys were used to make MCAO by the microcatheter method. MR scans were performed at 0 h (1 h after MCAO), 1 h, 3 h, 6 h, 12 h, 24 h, and 48 h after reperfusion. R2′ was calculated using quantitative T2 and T2∗ maps. Ischemic area was subdivided into infarcted core, IP, and oligemia. rR2′ was calculated respectively.

Results. Reversible MCAO model for 4/6 monkeys was made successfully. rR2′ values were significantly different at each time point, being highest in oligemia followed by IP and infarcted core (P<.05). With reperfusion time evolution, rR2′ in infarcted core showed a decreased trend: sharply decreased within 6 hours and maintained at 0 during 6–48 hours (P<.05). rR2′ values in IP and oligemia showed similar increased trend: sharply increased within 6 hours, maintained a plateau during 6–24 hours, and slightly increased until 48 hours.

Conclusion. BOLD-based R2′ MRI can be used to describe changes of cerebral oxygen extract in acute ischemic stroke, and it can provide additional information in detecting IP. The time evolution rR2′ in infarcted core, IP, and oligemia is in accordance with the underlying pathophysiology.

1. Introduction

The mismatch model between MRI perfusion-weighted imaging (PWI) and diffusion-weighted imaging (DWI) has been suggested to provide an estimation of real ischemic penumbra (IP) and is used as referring criteria to select patients who are within the first 6 hours after stroke onset and might benefit from thrombolysis [1–3]. However, recent studies indicated that the mismatch region is only a weak approximation of IP [4]. PWI cannot reliably discriminate between benign oligemia and penumbra, and DWI lesions do not always indicate the irreversible core of infarction [5]. It is well established that PET is a more accurate imaging modality to detect IP, which has decreased cerebral blood flow (CBF) but elevated oxygen extraction fraction (OEF), reflecting a compensatory mechanism in still-viable but threatened tissue [4–6]. However, PET is not suitable for emergency imaging study of acute cerebral infarction because of its limited availability and cost effectiveness.

The deoxyhemoglobin (deoxy-Hb) in cerebral capillaries and veins is an indicator of OEF, which can be visualized by T2-and T2*-based blood-oxygen-level-dependant (BOLD) MRI [7]. The potential for detecting focally increased OEF as reduced T2* signal was first reported by Roussel et al. [8] in rats middle cerebral artery occlusion (MCAO) model in 1995. Tamura et al. [7] also reported signal loss on T2* WI obtained before arrival of contrast medium on PWI in patients with acute ischemic stroke, potentially reflecting a high OEF in 2002. Similar MRI findings were also reported by Wardlaw and Von Heijne [9] in 2006 and Morita et al. [10] in 2008, respectively. However, the relationship between T2* and OEF is still controversial. In a small series of 5 patients, Donswijk et al. [11] found, despite of all patients showing increased OEF on PET, no significant relationship between OEF and T2* signal obtained before arrival of contrast medium either within or between subjects. This may be due to the fact that T2* is sensitive to large variations of the static magnetic field (ΔB0). R2′ is referred to as reversible
transverse relaxation rate, a derivational parameter from \( T_2^* \) and \( T_2 \) (\( 1/T_2^* = 1/T_2^+ - 1/T_2, \ R_2^* = 1/T_2^+ \), \( R_2 = 1/T_2, \ R_2' = 1/T_2^' \)) [12, 13]. The use of \( R_2^* \) is reasonable because it is isolated from the influences of deoxy-Hb on \( T_2^* \) values and water content on \( T_2 \) values. Several studies have used \( T_2' \) to predict lesion growth in acute ischemic stroke and demonstrated that it is sensitive to local content of deoxy-Hb [12, 14, 15].

The purposes of this study were to investigate: (1) the value of \( R_2' \) in detecting IP in a monkey model of reversible MCAO; (2) time evolution of relative \( R_2' \) (r\( R_2' \)) in infarcted core, IP, and oligemia.

2. Materials and Methods

2.1. Monkey Model of Reversible MCAO. This study was approved by the experimental animal ethical committee of our university. Six male monkeys (Macaca mulatta) with a mean weight of 8.6 kg (range: from 8 to 10 kg) were supplied by Experimental Animal Center of Beijin Military Medical Academy. All animals were fasted for 12–15 hours before experiment. Anesthesia was induced by ketamine (0.25 mg/kg intramuscular) and atropine (8–10 mg/kg intramuscular) followed by 100 mg ketamine and 50 mg diazepam in 500 mL NS intravenous drip. Physiological parameters such as respiration, heart rate, body temperature, and blood pressure were monitored.

The MCAO model was established by an interventional microcatheter method as described by De Grespigny et al. [16]. A one-inch incision was made in right femoral groove, microcatheter method as described by De Grespigny et al. were monitored.

The 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 experiment. Anesthesia was induced by ketamine (0.25 mg/kg intramuscular) and atropine (8–10 mg/kg intramuscular) followed by 100 mg ketamine and 50 mg diazepam in 500 mL NS intravenous drip. Physiological parameters such as respiration, heart rate, body temperature, and blood pressure were monitored.

2.2. Imaging Study. MRI scan was performed on a 1.5 T superconductive scanner (GE Twinspeed Infinity with Excite I) with birdcage head 8-channel phase array coil. All animals were scanned at 0 hour (1 hour after MCAO) and 1, 3, 6, 12, 24, and 48 hours after reperfusion. Slice thickness was 5 mm and FOV was 15 cm x 15 cm. Scanning protocol included, (1) T2WI (fast recovery FSE, TR 9002 ms, TE 124 ms, NEX 2, matrix 256 x 256). (2) DWI (SE-EPI, b values of 0 and 1000 s/mm²), TR 3000 ms, TE 112.6 ms, NEX 2, matrix 128 x 128) (3) Quantitative T2-map (qT2) (4-echo SE, TR 1200 ms, TE 25, 50, 75, 100 ms; flip angle 150°, matrix 256 x 256) and Quantitative T2*-map (qT2*) (8-echo GRE, TR 150 ms, TE 3.6, 9.4, 15.3, 21.2, 27.1, 33.0, 37.5, 43.2 ms, flip angle 24°, matrix 256 x 256) (4) PWI (single-shot gradient recall echo planner imaging T2* WI, TR 2000 ms, TE 80 ms, flip of angle 90°, matrix 128 x 128, NEX 1, Gd-DTPA 0.1 mmol/kg, inject rate 2 mL/s).

\( \text{qT2-map and qT2*-map were calculated according to the following formula:} \ SI(t) = SI_0 e^{-t/T2} \text{ and} \ SI(t) = SI_0 e^{-t/T2*}. \text{ qT2-map is linked to qT2*-map, resulting in} \ R_2' \text{-based BOLD imaging as follows:} \ 1/T_2' = 1/T_2^* - 1/T_2, \ R_2^* = 1/T_2^*, \ R_2 = 1/T_2, \ R_2' = 1/T_2' \ R_2^* \text{ is total transverse relaxation rate;} \ R_2 \text{ is irreversible transverse relaxation rate;} \ R_2' \text{ is reversible transverse relaxation rate with the unit of Hz [12, 13].} \)

2.3. Postmortem and Pathological Studies. At 48 hours after reperfusion, all animals were killed by an overdose of pentobarbital just after MRI scan. The brains were put into a −20°C refrigerator for an hour to make the specimens hardened. The brains were sliced into 5-mm axial slabs by using a home-made instrument and stained with 2,3,5-triphenyltetrazolium chloride (TTC) in 37°C water bath for 30 minutes. The stained sections were recorded with a digital camera.

2.4. Imaging Analysis and Statistics. Lesion volumes were calculated on DWI at 0 hour (0-DWII), T2WI at 48 hours after reperfusion (48-T2WI)- and TTC-stained sections by integrating products of lesion area and slice thickness on each lesion-containing slice. Lesion volume percentage (LVP) was calculated according to formula: \( \text{LVP = lesion volumes/bilateral hemispheres volumes x 100\%.} \)

Ischemic brain tissues were classified into 3 subtypes: infarcted core (hyperintensive areas on both 0-DWI and 48-T2WI), IP (hyperintensive areas on 0-DWI but iso on 48-T2WI), oligemia (delayed mean transit time areas on PWI at 0 hour [0-MTT] but iso on both 0-DWI and 48-T2WI) (Figures 1, 2, 3, and 4).

Imaging registration was performed on an off-line workstation. The platform was Matlab 7.0, Windows XP SP2 with 3.0 GHz Intel Pentium 4 CPU and 2 G RAM. 0-DWI, 0-MTT, and 48-T2WI were coregistered to R2’ images of each time point using a free form deformation method [17].

5 region-of-interests (ROIs) were put into each subtype brain tissue, respectively. Each ROI was 10 mm² in size. Relative R2’ (rR2’) was calculated according to the ratio of lesion and its minor area on contralateral healthy hemisphere.

Statistic analysis was performed with SPSS15.0 software. Paired t-test and correlation analysis were used for comparison of LVP. rR2’ values in three-subtype brain regions at
3. Results

3.1. MCAO Model. MCAO model was successful in all 6 monkeys. 2 monkeys were dead at 2 and 14 hours after reperfusion because of large area of hemorrhagic infarction, respectively, and their data were not included in this study.

3.2. LVP. 0 h DWI demonstrated high-signal ischemic lesion and partial lesion can recover after reperfusion (Figures 5, 6, and 7), which was confirmed by TTC staining (Figure 8).

LVP on 0-DWI, 48-T2WI- and TTC-stained sections were summarized in Table 1. LVP on 48-T2WI was decreased compared with that on 0-DWI (t = 6.472, P < .05, ranged: 23% ~35%, average 28%). There was no statistic significant between LVP on 48-T2WI and that on TTC (t = 0.15, P > .05), which showed significant positive correlation (r = 0.98, P < .05).

Table 1: LVP on DWI at 0 hour, T2WI at 48 hours after reperfusion and TTC.

<table>
<thead>
<tr>
<th>Animal number</th>
<th>DWI (%)</th>
<th>T2WI (%)</th>
<th>TTC (%)</th>
</tr>
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<tbody>
<tr>
<td>1</td>
<td>10.77</td>
<td>8.08</td>
<td>8.19</td>
</tr>
<tr>
<td>2</td>
<td>12.78</td>
<td>8.31</td>
<td>8.45</td>
</tr>
<tr>
<td>3</td>
<td>12.26</td>
<td>8.77</td>
<td>8.68</td>
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<tr>
<td>5</td>
<td>9.68</td>
<td>7.48</td>
<td>7.27</td>
</tr>
</tbody>
</table>

3.3. rR2′ in Different Ischemic Brain Tissue. On 0 hour R2′ imaging, IP and oligemia showed hyperintensity compared to infarcted core. After reperfusion, the signal intensity on each time point were compared by ANOVA. Time evolution of rR2′ in each subtype brain region was also compared by ANOVA. Statistic significance was P < .05.

rR2′ imaging was increased in IP and oligemia but decreased in infarcted core with time evolution (Figures 9, 10, and 11). rR2′ values of infarcted core, IP, and oligemia at each time point after reperfusion were summarized in Table 2. rR2′ values were significantly different at each time point, being highest in oligemia followed by IP and infarcted core (P < .05).

3.4. Time Evolution of rR2′ in Different Ischemic Brain Tissue. Respective time evolution of rR2′ values in infarcted core, IP, and oligemia was summarized in Table 3. The infarcted core showed a decreased trend after reperfusion (P < .05): the maximum decrease happened within the first 6 hours and then maintained as low level as 0 during 6–48 hours. While rR2′ values in IP and oligemia showed similar increased
4. Discussion

4.1. Monkey Model of MCAO. As described by De Crespigny et al. [16], we successfully made monkey model of reversible MCAO by an interventional method in 4/6 monkeys and performed dynamic MRI scans for them. No artifact was found since we removed the microguide wire during MR scan. Unfortunately 2 monkeys died of large area of hemorrhagic infarction in this study. This may due to the fact that we used 250,000 units urokinase to ensure effective reperfusion after the micocatheter was removed.

Although many neuroprotective drugs have proven successful in animal (rodent) models, all have failed during clinical trials. A common reason for this is the lack of demonstrated efficacy of these drugs in larger animals such as cats or monkeys. The subhuman monkeys have similarity in cerebrovascular anatomy, neurochemistry, and immunology, which make them suitable for testing new drug therapies and endovascular instruments [16].

4.2. BOLD-Based $R_2'$ MRI in Different Ischemic Brain Tissues.

The delineation of the “penumbra” is of particular interest in acute stroke imaging. Since the “mismatch concept” applying PWI and DWI appears to be an oversimplification of the underlying electrophysiological tissue status, an additional parameter reflecting the metabolic state of brain tissue will improve the ability to describe the penumbra [12]. BOLD MR T2‘ imaging is thought to reflect the response of tissue metabolism to that perfusion deficit with increased local deoxy-Hb used as an indicator of increased OEF. Several studies confirmed the additional values of T2‘ map in predicting the real penumbra [12, 14, 15]. In this study, we used $R_2'$ to describe changes of local deoxy-Hb. Increased local deoxy-Hb leads to hyperintensity on $R_2'$ images, while decreased local deoxy-Hb leads to hypointensity on $R_2'$ images. This makes it easier to visually notice changes of local deoxy-Hb.

Our study demonstrated that the values of $R_2'$ were increased in IP and oligemia and decreased in infarcted core with time evolution after reperfusion. Thus, the hyperintensity in $R_2'$ images may represent increased deoxy-Hb as an indicator of increased OEF, which can increase in IP and oligemia with ongoing CBF impairment. The infarcted core is unsalvable tissue which has lost the compensative ability of...
Table 2: Comparison of rR2′ (\(\overline{x} \pm s\)) in infarcted core, IP, and oligemia at each time point after reperfusion.

<table>
<thead>
<tr>
<th>Subtype</th>
<th>0 h</th>
<th>1 h</th>
<th>3 h</th>
<th>6 h</th>
<th>12 h</th>
<th>24 h</th>
<th>48 h</th>
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</thead>
<tbody>
<tr>
<td>Infarcted core</td>
<td>1.129</td>
<td>0.108</td>
<td>0.668</td>
<td>0.082</td>
<td>0.536</td>
<td>0.075</td>
<td>0.259</td>
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<tr>
<td>IP</td>
<td>1.329</td>
<td>0.081</td>
<td>1.237</td>
<td>0.072</td>
<td>1.453</td>
<td>0.081</td>
<td>2.435</td>
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<tr>
<td>Oligemia</td>
<td>1.584</td>
<td>0.103</td>
<td>1.435</td>
<td>0.066</td>
<td>1.770</td>
<td>0.141</td>
<td>2.957</td>
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<tr>
<td>(F)</td>
<td>36.19</td>
<td>134.09</td>
<td>256.30</td>
<td>803.25</td>
<td>743.74</td>
<td>1236.26</td>
<td>557.02</td>
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<td>(P)</td>
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Table 3: Time evolution of rR2′ in infarcted core, IP, and oligemia after reperfusion (compared by ANOVA).

<table>
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<tr>
<th>Reperfusion time</th>
<th>Compared time</th>
<th>IP</th>
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<th>Oligemia</th>
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\(* P < .05, — P > .05.\)

Figure 9: 0 h R2′ map shows that the signal of IP (black arrow) is higher than that of infarcted core (white arrow).

An interesting finding demonstrated by Geisler et al. [12] was the infarcted core showed even shorter T2′ than lesion growth (operationally defined IP, −15.7% versus −10.5%) although there was no statistically significant difference. That result was not as they have expected: a strongly reduced O2 utilization, a clearly lowered OEF, a decreased deoxy-Hb, and consecutively increased T2′ but also include a considerable portion of IP which showed decreased T2′, and may thus account for the T2′ shortening [12].

Although both IP and oligemia showed similar increased trend in this study, values of rR2′ in oligemia were higher.

increased OEF. Thus, the infarcted core demonstrated consecutive hypointensity in R2′ images even after reperfusion. PET study has showed 50% increasement of OEF in IP [18].
than that in IP at every time point after reperfusion. There are two reasons for this. Firstly, CBF in oligemia is less impaired than those in IP [21], which means that the oxygen supply in oligemia is more sufficient. Secondly, the compensative ability of oligemia is stronger than that of IP, which leads to more increased OEF and increased generation of deoxy-Hb. The less decreased CBF and more increased OEF are the real reasons that oligemia can escape from ischemic injury and never develop to infarction. Certainly, we can establish a $rR2'$ threshold to distinguish IP and oligemia. Considering the limited number of our animal models, we would leave this job for further study.

4.3. Time Evolution of $rR2'$ in Different Ischemic Brain Tissues.

Both PET and BOLD MRI studies had showed increased OEF in acute ischemic stroke [4–6, 12, 14, 15]. But the duration of increased OEF is still unknown. According to the results of our study, time evolution of $rR2$ in IP and oligemia can be classified into 3 stages: sharply increased within first 6 hours, maintained a high level plateau during 6–24 hours, and then slightly increased until 48 hours. While the infarcted core had 2 stages: sharply decreased within first 6 hours and maintained as low level as 0 from 6 to 48 hours. 6 hours is also the time window for venous thrombolytic therapy in acute ischemic stroke and is widely used in many stroke studies [1, 4, 6, 12, 14, 18, 22]. Early recanalization is the first and most important step to save IP. The most obvious hemodynamic changes may happen during the first 6 hours after early reperfusion [21]. We presume quick recovered CBF may encourage IP to increase OEF excessively, generate more deoxy-Hb, and lead to sharply increased $rR2'$. With the gradual recovery of CBF from 6 to 24 hours, IP may still be in moderate anoxic status and OEF may still increase at a lower speed. The plateau during 6–24 hours may represent the balance state between recovering CBF and elevated OEF. At 48 hours CBF may have already completely recovered but OEF may still be maintained at a higher level, which leads to slightly increased $rR2'$. That is to say, increased OEF may last at least 48 hours after early reperfusion. A previous study confirmed that extended duration of increased OEF will benefit brain tissue at risk [22]. 48 hours duration of increased deoxy-Hb disclosed by our study may be helpful to extend therapy time window of ischemic stroke.

For the infarcted core, the sharply decreased $rR2'$ within the first 6 hours may be due to the fact that no generation of deoxy-Hb is found even after early reperfusion. With the quick recovery of CBF, the previously produced deoxy-Hb is removed and $rR2'$ maintains as low level as 0 after the first 6 hours.

There are some methodological limitations for $T2'$ or $R2'$ images. Firstly, $T2'$ is sensitive to large variations of the static magnetic field ($\Delta B0$). This may affect the accuracy of measured $T2'$ or $R2'$. This problem can be solved by using side-to-side comparison which reveals more valuable relative results. Secondly, the concentration of deoxy-Hb in a single voxel is determined mainly by 2 factors: changes in the local generation of deoxy-Hb inside the voxel and changes in the transport of deoxy-Hb into or out of the voxel,

Figure 10: 1 h R2' map shows low signal of irreversible infarcted core (white arrow), slightly high signal of reversible IP (black arrow).

Figure 11: 48 h R2' map shows significant low signal of infarcted core (white arrow), high signal of IP (black arrow) (Figures 9, 10, and 11 is same animal as Figures 5, 6, and 7).

Figure 12: Reperfusion time evolution of $rR2'$ shows decreased trend in infarcted core but increased trend in IP and oligemia.
which is closely correlated with CBF [12]. So information of CBF should be imposed on T2’ or R2’ images. A study carried by Jensen et al. [15] showed that there was a moderate correlation between rCBF derived from PWI and qT2’. qT2’-maps could be used to explore cerebral perfusion without the application of contrast agent or radiation. Although BOLD-based MR quantitative OEF measurement has been published and demonstrated good correlation with the existing OEF values [23, 24]. The clinical value of this technology is still unknown. Further studies should be focused on the improvement of accuracy of cerebral oxygen extraction by using BOLD MR.

5. Conclusions

BOLD-based R2’ MRI can be used to describe changes of cerebral oxygen extract in acute ischemic stroke, and it can provide additional information in detecting IP. The time evolution rR2’ in infarcted core, IP, and oligemia is in accordance with the underlying pathophysiology. With improvement in the accuracy, this technique would add more valuable information for patients selection for thrombolytic therapy.

Acknowledgments

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