Abstract. Background/Aim: Middle cerebral artery occlusion (MCAO) in rodents is an essential animal model for research focusing on ischemic stroke. To date, several kinds of surgical methods for MCAO have been developed and the craniotomy method has the advantage of direct visualization of the middle cerebral artery (MCA). MCAO at a more proximal site produces better surgical results, but it is a more invasive technique. The aim of this study was to evolve the surgical technique for simulating ischemic cerebral cortex injury in rats. Materials and Methods: To approach proximal MCA with a less invasive procedure, a modified surgical technique for MCAO in rats was developed. Besides, rats receiving the modified and conventional method were compared with regard to infarct volume and by behavioral tests. Results: Following craniotomy, we proposed that the inferior edge of the craniotomy should be enlarged with fine forceps. This modified surgical method induces larger infarct volume, significant behavioral impairment and can induce ischemic stroke. Additionally, it does not significantly increase the operation time, and has produced no obvious complications. Conclusion: This modified surgical technique may serve as a practical method for performing MCAO.

Stroke is a common disease that can have severe sequelae (1). About 80% of strokes are ischemic, and the most common infarction site is the middle cerebral artery (MCA) (2, 3). To facilitate progress in stroke research, translational research in animal models is essential in order to transfer basic research findings to the clinic. Researchers regularly choose the rodent model to create middle cerebral artery occlusion (MCAO) to simulate ischemic stroke (4-6). There exist several types of methods to induce MCAO, including intraluminal filament occlusion, craniotomy and direct MCA occlusion, and Endothelin-1-induced MCAO (7, 8).

Among these methods, craniotomy is a common method, that has the advantage of allowing direct visualization of MCA through the surgical field (9). After the MCA has been exposed, it can be occluded by coagulation, transection, ligation or clips (7). However, the precise location of MCA occlusion is a critical issue. One study that used craniotomy in rats for MCAO revealed that occlusion of a more proximal part of the MCA led to larger infarct size (10). However, removing the cranial bone over the proximal MCA during craniotomy may be impeded by the zygomatic arch. If operators want to expose the most proximal part of the MCA, the zygomatic arch may be resected to provide adequate space (11, 12). However, this process may take time or affect the eating ability of rats. Other authors have exposed proximal MCA by displacing the brain but the large displacement of the brain may affect physiological functions (10). Intraluminal filament occlusion is another MCAO model that can mimic the clinical pathophysiology of stroke;
however, this method will also obstruct anterior cerebral artery partially and produce striatum infarction (5). Thus, if researchers want to explore pure cortex ischemic injury without striatum invasion, it is not suitable to use intraluminal filament occlusion as an MCAO model. 

Other than producing large and stable infarct size, a competent MCAO model is also expected to cause obvious neurological deficits thus simulating ischemic stroke. After ischemic brain injury, inflammatory cytokines such as tumor necrosis factor (TNF-α), interleukin (IL)-1β, and IL-6 are produced in the ischemic brain tissue and this neuroinflammation is a major mechanism in ischemic brain injury (13). The change of these cytokines were verified in vivo and in vitro. Thus, a satisfactory MCAO model should also produce these major changes in inflammatory cytokines.

In this study, a modified surgical method to expose proximal MCA was developed. Following craniotomy, the inferior edge of the craniotomy was enlarged with fine forceps. The method was easy to perform without significantly extending the operation time. Furthermore, this method did not lead to obvious complications such as reduced eating ability or mortality. Finally, the method was able to recapitulate significant effects of stroke, such as large and stable infarct volume, impaired behavior performance and significant production of inflammatory cytokines. Furthermore, the results of proximal and distal MCAO were also compared.

Materials and Methods

Animals. All animal experiments were approved by the Institutional Animal Care and Utilization Committee (IACUC) of National Yang-Ming University, Taipei, Taiwan (approval ID: 1050315). Ten-week-old male Wistar Kyoto (WKY) rats weighing 200-300 g each were purchased from BioLASCO (Taipei, Taiwan). The rats were housed in the Laboratory Animal Center in National Yang-Ming University with a 12-h light/dark cycle at a constant temperature (25±2˚C), and with unrestricted access to water and food.

Modified surgical method of MCAO. In order to improve the efficiency of MCAO, a modified surgical method was developed. Rats were anesthetized with an intraperitoneal injection of Zoletil (30 mg/kg) and Xylazine (8 mg/kg). A vertical incision was made on the midline of the neck, and the bilateral common carotid arteries (CCAs) were identified. After retracting the temporal muscle, a craniotomy was performed on the right temporal bone using a cutting burr (Figure 1a and b). During this period, the inferior edge of the hole was lowered as much as possible, until interrupted by the zygomatic arch. Then Dumont forceps (Roboz, Gaithersburg, MD, USA) were used to hold the residual bone on the outer surface of cranial bone, and another tip on the outer surface of cranial bone (Figure 1c). After holding it with about 1mm contact depth, the bone was pulled out piecemeal with a lateral and superior direction until the olfactory tract was exposed (Figure 1d). Then the dura matter was removed, and the MCA was permanently occluded by suture ligation at the level of the olfactory tract. Subsequently, bilateral CCA occlusion was performed by vessel clamp temporarily for 90 min.

Another rat group was subjected to MCAO for purposes of comparison, using the conventional method. In this group, rats underwent the above procedure, but forceps were not used to remove the residual bone on the inferior edge of the hole. Thus, MCA was exposed to the level above the inferior cerebral vein and the distal MCA was permanently occluded by suture ligation above the inferior cerebral vein. Bilateral CCA occlusion was also performed temporarily for 90 min. The sham group received identical anesthesia and surgical procedure without MCA and CCA occlusion. Rats that died during or after surgery were excluded for further experiment (n=7).

Animal Behavioral Test. The rotarod test was utilized to evaluate motor coordination and balance alterations. Before MCAO was applied, rats were trained for running and balancing on the rotating drum (Rotamex-5, Columbus Instruments, Columbus, OH, USA) for 3 days. The rats deemed stable and that performed well on rotarod were assigned to groups that randomly received modified and conventional MCAO. At 1 (n=13/group) and 3 (n=7/group) days after MCAO, the rotarod was accelerated from 4 rpm to 40 rpm, with an acceleration of 1 rpm/10 sec. The latency before falling off the drum was recorded. Every rat received three trials for each training and testing day and the interval between each trial was 10 min. Successful trials after MCAO were defined as less than 75% running time on rod at 1 day after surgery, compared with preoperative performance.

Quantification of Infarct Volume. At 1 (n=8/group) and 3 (n=4/group) days after MCAO, rats were anesthetized by intraperitoneal injection of Zoletil and Xylazine and transcardially perfused with phosphate-buffered saline (PBS). Their brains were expeditiously removed from the skull to determine infarction size, and then sliced into 2-mm-thick coronal sections. Then, they were stained with 2%, 3,5- triphenyltetrazolium chloride (TTC; Sigma, St. Louis, MO, USA) solution for 15 min at room temperature without light, and then fixed with 4% paraformaldehyde. The unstained area represented the infarcted region, and the non-infarcted region revealed as red color on TTC-stained sections. To eliminate the effect of brain edema, the infarcted region was determined by subtracting the non-infarcted region in the ipsilateral hemisphere from that in the contralateral hemisphere. Infarct volumes were then calculated as the sum of the infarcted areas on each brain slice by using Image J software (National Institutes of Health, Bethesda, MD, USA).

Measurement of inflammatory cytokines. At 1 (n=4/group) day after MCAO, rats were anesthetized by intraperitoneal injection of Zoletil and Xylazine and transcardially perfused with PBS. Their brains were removed and stored in –80˚C ultra-low temperature freezer. Then, the cerebral cortex of MCAO rats and sham group were homogenized and the levels of inflammatory cytokines (IL-1β and IL-6) were measured using commercially available ELISA kits (R&D, Minneapolis, MN, USA).

Statistical analysis. Data are expressed as mean±the standard error of the mean (SEM). Categorical data were compared using chi-square test. Independent samples t-test was used to analyze the difference between two groups. All statistical analyses were
performed using the SPSS 24.0 software package (SPSS, Inc., Chicago, IL, USA). Differences were considered significant at \( p<0.05 \).

**Results**

**Operation time and mortality rate of the surgical procedures.** The mean operation time from the point of skin incision to completion of a 90-minute CCA occlusion of rats receiving the modified and conventional method was 130.2 min and 121.4 min, respectively. Following MCAO, the mortality rates from the modified and conventional surgical methods were 9.4% and 13.3%, respectively (\( p=0.622 \)). With the exception of the neurological deficits induced by MCAO, none of the rats had other obvious complications including eating difficulties.

**The rats that received the modified surgical technique performed poorer in the rotarod test.** Before MCAO, the rats receiving the modified and conventional surgical methods

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**Figure 1.** The procedures of the modified surgical method for middle cerebral artery occlusion (MCAO). (A) After skin incision and temporal muscle retraction, craniotomy contour was created using a burr tool. (B) Craniotomy and exposure of the underlying brain tissue. (C) Removal of the residual bone on the inferior edge of the craniotomy (arrow) using fine forceps. (D) Exposure of the middle cerebral artery at the level of olfactory branch (arrow) and preparation for MCAO.
had similar running times on the rod (126.1 sec vs. 125.6 sec, \( p = 0.930 \)) (Figure 2a). At 1 day after MCAO, the group that received the modified surgical procedure had less running time on the rod, compared with the group that received the conventional surgical procedure (51.6 sec vs. 90.5 sec, \( p = 0.009 \)) (Figure 2a). At 3 days after MCAO, rats that received the modified method still stayed less time on the rod, compared with rats that received the conventional method (49.9 sec vs. 96.9 sec, \( p = 0.071 \)) (Figure 2a). The group that received the modified surgical procedure had a success rate of 76.9\%, whereas the group that received the conventional surgical procedure had a success rate of only 46.2\% (Figure 2b).

The modified surgical technique induced larger infarct volume. The ischemic region revealed as a pale color by TTC stain and involved only the cortex (Figure 3a and b). At 1 day after MCAO, the infarct volume of the rats that received the modified surgical method was 189.5 mm\(^3\), larger than the 115.4 mm\(^3\) of the group that received the conventional surgical method (\( p = 0.001 \)) (Figure 3c). At 3 days after surgery, the infarct volumes in the groups operated with the modified and conventional methods were 144.6 mm\(^3\) and 102.3 mm\(^3\), respectively (\( p = 0.004 \)) (Figure 3c). With time, the infarct size became smaller in the group that received the modified surgical method (\( p = 0.021 \)) (Figure 3c).

The modified surgical technique increased the levels of inflammatory cytokines. At 1 day after MCAO, the mean concentration of IL-1\( \beta \) in the ischemic cortex was 319.2 pg/ml, higher than the 44.3 pg/ml of the sham group (\( p = 0.039 \)) (Figure 4). The mean concentration of IL-6 in the ischemic cortex was 1,024.1 pg/ml and in the sham group 70.9 pg/ml (\( p < 0.001 \)) (Figure 4).

Discussion

In the present study an efficient MCAO method via craniotomy in order to simulate ischemic cerebral cortex injury in rats is presented. Among the MCAO methods, intraluminal occlusion is a common method that requires skilled technique and expensive laser Doppler flowmetry to detect changes in MCA blood flow (14). It does not allow for direct observation of MCA. Besides, intraluminal occlusion usually induces infarction extending to striatum, so this occlusion method may not be suitable for studying exclusive cortex infarction. Alternatively, MCAO by way of craniotomy does not necessitate the use of expensive equipment, and can allow direct visualization of MCA. However, the site of MCAO significantly affects the results (10).

To induce larger infarct volume, the MCA must be occluded at the proximal site, such as the level of olfactory...
branch (10). However, if a burr tool is used to perform craniotomy, the burr shaft will be impeded by the zygomatic arch. Even if a burr with a specific angle is utilized, the procedure will still be impeded due to the width of the burr head. Removing the zygomatic arch is another option available to occlude the MCA at the proximal site. However, this option will likely extend the operation time, and may interfere with the eating ability of rodents.

Figure 3. The modified surgical method produced larger infarct volume at 1 and 3 days after middle cerebral artery occlusion (MCAO). Representative, TTC stained brain slices from rats that received sham-operation, the modified surgical method or the conventional surgical method at (A) 1 day and (B) 3 days after MCAO. The pale region indicates the infarct area. (C) Quantification of infarct volume of rats that received the modified or conventional surgical method on 1 day and 3 days after MCAO. The sham-operated rats showed no infarction. n=8/group in 1 day and n=4/group in 3 days after MCAO. Data are presented as mean±SEM. **p<0.05.

Figure 4. The modified surgical method produced higher levels of inflammatory cytokines in the ischemic cortex compared to the sham operation at 1 day after middle cerebral artery occlusion (MCAO). (A) The concentration of IL-1β in the cerebral cortex of MCAO and sham groups. (B) The concentration of IL-6 in the cerebral cortex of MCAO and sham groups, n=4/group. Data are presented as mean±SEM. *p<0.05, ***p<0.001.
Thus, we have developed this novel and easy technique to improve the efficiency of MCAO by means of craniotomy. The key point in this modified procedure is to use fine forceps to remove the residual inferior cranial bone piecemeal, working in a lateral and superior direction. If a medial direction force is misapplied, the brain may be injured and the MCA could be impacted, potentially causing a hemorrhagic stroke. If an inferior direction force is misdirected, the residual bone cannot likely be efficiently removed. We suggest that forceps should be used to hold the bone with a 1 mm contact depth. If contact is shorter than this depth, the residual bone is difficult to remove; if longer than this depth, brain tissue injury may occur. This modified method can produce large infarct volume; however, the infarct area is confined to the cerebral cortex. Thus, this method may not be suitable when focusing on other areas such as striatum.

It is generally understood that bilateral CCA occlusion can increase the infarct volume due to additional block of the collateral circulation of the brain (15). If occluding the unilateral or bilateral CCA permanently, the mortality rate increases significantly (15). Chen et al. have reported an elevated mortality rate of 60% in rats receiving right MCA and bilateral CCA permanent occlusion (15). Thus, in this modified surgical procedure, temporal bilateral CCA occlusion was applied. This can increase the infarct volume without increasing the mortality rate.

Current studies focusing on the methodology of MCAO usually utilize infarct size and neurological deficits to evaluate if the MCAO model is successful. Changes in the cytokine levels are seldomly examined. Thus, to reinforce the evidence of a successful MCAO model, production of inflammatory cytokines was also measured and proved that our modified method increased the secretion of IL-1β and IL-6 in the ischemic cerebral cortex.

In conclusion, this study describes a modified surgical method to perform MCAO by way of craniotomy, with satisfactory results. This method simulates ischemic cerebral cortex injury, and produces large infarct volume and impaired motor and balance function. Besides, inflammatory cytokines were significantly increased in the ischemic region. Furthermore, further evidence indicated that this occlusion of the more proximal part of MCA by the modified surgical procedure can have a more severe stroke impact compared to the conventional method.

Conflicts of Interest

The Authors declare that they have no competing interests in regard to this study.

Authors’ Contributions

Conceptualization: Chien-Fu Yeh, Tung-Yueh Chuang, Ming-Ying Lan and Yung-Yang Lin; Data curation: Chien-Fu Yeh, Tung-Yueh Chuang, Yu-Wen Hung, Ching-Han Tsai and Hao-Xiang Huang; Supervision: Ming-Ying Lan and Yung-Yang Lin; Writing – original draft, Chien-Fu Yeh; Writing – review & editing: Chien-Fu Yeh, Tung-Yueh Chuang, Ming-Ying Lan and Yung-Yang Lin.

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