A Nomogram Model to Predict Malignant Cerebral Edema in Ischemic Stroke Patients Treated with Endovascular Thrombectomy: An Observational Study

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Purpose: Malignant cerebral edema (MCE) in patients undergoing endovascular thrombectomy (EVT) is not uncommon and can reduce the benefit of EVT. We aimed to develop a nomogram model to predict the risk of MCE in ischemic stroke patients after EVT.

Patients and Methods: We retrospectively collected patients treated with EVT caused by anterior circulation large vessel occlusion stroke at two comprehensive stroke centers. MCE was defined as midline shift >5 mm at the septum pellucida or pinal gland with obliteration of the basal cisterns or the need for early decompressive hemicraniectomy. A multivariate logistic model was utilized to construct the best-fit nomogram model. The discrimination and calibration of the nomogram were estimated using the area under the receiver operating characteristic curve (AUC-ROC) and Hosmer–Lemeshow test.

Results: A total of 370 patients (mean age, 67.2±11.9 years; male, 56.8%) were enrolled in the final analysis. Among them, 71 (19.2%) patients experienced MCE after EVT treatment. After adjustment for potential confounders, age, baseline National Institutes of Health Stroke Scale score, collateral circulation, fast blood glucose level and recanalization were independent predictors of MCE and were incorporated into the nomogram. The AUC-ROC value of the nomogram was 0.805 (95% confidence interval [CI]: 0.750–0.860). The Hosmer–Lemeshow goodness-of-fit test showed good calibration of the nomogram (P = 0.681).

Conclusion: The nomogram composed of age, baseline National Institutes of Health Stroke Scale score, blood glucose level, collateral circulation and recanalization may predict the probability of MCE in anterior circulation large vessel occlusion stroke patients treated with EVT.

Keywords: endovascular thrombectomy, edema, nomogram, stroke

Introduction
Randomized controlled trials have validated the efficacy and safety of endovascular thrombectomy (EVT) in patients with anterior circulation large vessel occlusion stroke (LVOS).1–5 Moreover, more patients could benefit from EVT based on imaging screening.6,7 However, approximately 45% of patients with LVOS are still unable to perform functional recovery after EVT.8,9 Recent studies have shown that malignant cerebral edema (MCE) after EVT is not uncommon and might reduce the benefit-risk ratio of EVT.10–12 MCE is one of the most serious complications of acute ischemic stroke (AIS), and the mortality rate in MCE
patients is nearly 80%.13 Although treatments of MCE are limited, a pooled analysis of randomized-controlled trials shows that early decompressive hemicraniection could decrease mortality and increase the chance of a favorable functional outcome.14 Therefore, the early prediction of MCE might bring more benefits for patients with AIS.

A recent systematic review indicated that the major determinants for MCE after stroke are younger age, high National Institutes of Health Stroke Scale (NIHSS) score, larger parenchymal hypoattenuation and the degree of reperfusion.15 Additionally, several studies in patients with intravenous thrombolysis (IVT) show that baseline blood glucose, onset-to-treatment time (OTT) and hyperdense artery sign were independent predictors of MCE.16,17 However, data on the prediction of MCE in patients treated with EVT are relatively scarce. Hence, in this study, we aimed to develop a nomogram model based on demographic and clinical features to predict MCE in patients treated with EVT.

 Patients and Methods

Study Population

We retrospectively collected the demographic, clinical and radiographic data of patients with anterior circulation LVOS who underwent EVT at two comprehensive stroke centers (Jinling Hospital between January 2014 and December 2018 and Yijishan Hospital between July 2014 and February 2019). Patients were enrolled if they fulfilled the following inclusion criteria: (1) age ≥ 18 years; (2) time from stroke onset to puncture (OTP) ≤ 8 hours; (3) baseline NIHSS score ≥ 6, baseline Alberta Stroke Program Early CT score (ASPECT) ≥ 6, and prestroke modified Rankin Scale (mRS) score < 2; (4) the internal carotid artery (ICA) or proximal segment (M1 or M2) of the middle cerebral artery (MCA) occlusion confirmed by preoperative imaging. The treatment protocol and methods were published previously.9,10 We excluded patients with extensive subarachnoid hemorrhage and midline shift or decompressive hemicraniection because of postproce-
dural hemorrhagic transformation. The study was approved by the Ethical Committee of Jinling Hospital and Yijishan Hospital (2010NL-018). Due to its retro-
spective nature; patient consent was waived. All proce-
dures performed in studies involving human participants were in accordance with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The data of patients were stored securely and confidentially in Jingling and Yijishan Hospitals.

Malignant Cerebral Edema and Clinical Characteristics

According to the follow-up imaging within 3–5 days after EVT, MCE was defined as midline shift > 5 mm at the septum pellucidum or pineal gland with obliteration of the basal cisterns or the need for decompressive hemicraniection.11,18 All neuroimaging data were reviewed by 2 physicians who were blinded to the clinical data. In case of disagreement, consensus was sought by discussion with a third physician.

The baseline clinical data included demographics, medical history, baseline NIHSS and ASPECT score, the Trial of ORG 10,172 in Acute Stroke Treatment (TOAST) classification and laboratory measures. Procedural characteristics, including OTP, time from stroke onset to recanalization (OTR), occlusion site, types of the procedure, bridging and rescue treatments, were all recorded. Recanalization status after the procedure was evaluated based on the modified Thrombolysis in Cerebral Infarction (mTICI) grading system. Successful recanalization was defined as an mTICI score of 2b or 3. Collateral circulation was assessed according to retrograde contrast opacification of vessels within the occluded area. The collateral score was classified as follows: grade 0 was assigned if there was little or no significant reconstitution in the territory of the occluded vessel or less than one-third of the occluded territory, grade 1 was assigned if the collaterals reached less than two-thirds of the occluded territory, and grade 2 was assigned if the collateralization reached more than two-thirds of the territory or the prox-
imal main stem.19

The functional outcomes were assessed with a 90-day mRS score, and a favorable functional outcome was defined as mRS score ≤ 2.

Statistical Analysis

Continuous variables are presented as the mean±standard deviation (SD) or as the median (interquartile range, IQR). Categorical variables are presented as percentages. Continuous variables were analyzed using the Mann–Whitney U-test. Categorical variables were analyzed using the Chi-square test or Fisher’s exact test as appropriate. To construct the nomogram, multivariate regression analysis with Akaike information criterion was applied to select the significant predictors of MCE using a backward-
selection method that included variables with a P value < 0.2 in the univariate analysis. The nomogram was established according to the significant predictors in the multivariate regression analysis. Regression coefficients and odds ratios (ORs) with two-sided 95% confidence intervals (CIs) for each variable included in the model were finally calculated. The collinearity of variables was evaluated by variation inflation factors.

The discriminative ability of the new nomogram was assessed by the area under the receiver operating characteristic curve (AUC-ROC). In addition, bootstrapping was used for internal validation. Calibration of the nomogram was assessed by a calibration plot and Hosmer-Lemeshow test. The calibration plot is a graphic showing the relationship between the observed outcome frequencies and the predicted probabilities. In a well-calibrated model, the predictions should fall on a 45-degree diagonal line. Statistical analyses were computed using R version 3.0 (R Foundation, Vienna, Austria) and SPSS 22.0 (IBM, Armonk, NY, USA). A two-tailed value of P < 0.05 was considered significant.

Results
In total, 370 patients (mean age, 67.2±11.9 years; male, 56.8%) were enrolled for the final analysis. The flow chart of the inclusion of the study population is displayed in Figure 1. Of the 370 patients treated with EVT, the median OTP was 247 minutes, and baseline NIHSS scores and ASPECT scores were 17.0 and 9.0, respectively. Among them, 313 (84.6%) patients received stent-retriever-first thrombectomy, 18 (4.9%) received aspiration-first, and 39 (10.5%) received angioplasty or stent-first procedure. Successful recanalization was achieved in 272 (73.5%) patients. During the follow-up at 3 months, 167 (45.1%) reached a favorable functional outcome (mRS, 0–2), and the overall mortality was 21.9%.

Predictors of Malignant Cerebral Edema
There were 71 (19.2%) patients who experienced MCE after EVT treatment. Table 1 shows the detailed demographic and clinical characteristics according to patients with and without MCE. In the univariate analysis, patients with MCE had a higher baseline NIHSS score (18.0 vs 16.0, P < 0.001) and a lower baseline ASPECT score (8 vs 9, P = 0.033). Furthermore, ICA-T occlusion (52.1% vs 24.7%, P < 0.001) and a lower degree of collateral circulation (grade 0, 42.3% vs 13.7%; grade 1, 43.7% vs 44.1%; grade 2, 14% vs 42.2%, P < 0.001) occurred more frequently in MCE patients. Additionally, patients with MCE received a higher rate of rescue treatment (28.2% vs 15.7%, P = 0.014) and spent a longer OTR (352 min vs 329 min, P = 0.015). However, the rate of successful recanalization (53.5% vs 78.3%, P < 0.001) was lower in patients with MCE. Additionally, a significantly higher level of fasting blood glucose (9.1 mmol/l vs 7.1 mmol/l, P < 0.001) and blood urea nitrogen (6.9 mmol/l vs 6.2 mmol/l, P = 0.036) was found in patients with MCE than in those without MCE.

After adjustment for potential confounders (variables with a P value < 0.2 in the univariate analysis), age (OR: 0.960, 95% CI: 0.934–0.987, P = 0.004), baseline NIHSS score (OR: 1.076, 95% CI: 1.016–1.140, P = 0.013), collateral circulation (grade 1 vs grade 0: OR: 0.360, 95% CI: 0.176–0.763, P = 0.005, grade 2 vs grade 0: OR: 0.127, 95% CI: 0.051–0.315, P < 0.001), fast blood glucose level (OR: 1.180, 95% CI: 1.086–1.281, P < 0.001) and recanalization status after the procedure (mTICI 2b–3 vs 0–2a, OR: 0.352, 95% CI: 0.185–0.669, P = 0.001) were independent predictors for MCE (Table 1). There was no significant statistical collinearity among the five independent predictors (Table S1).

A Nomogram Model for the Prediction of Malignant Cerebral Edema
The nomogram was developed by assigning a graphic initial score to each independent predictor with a point range from 0 to 100, which was then summed to generate...
Table 1 Comparison of Variables Between All Patients with and without Malignant Cerebral Edema

| Demographics | Total  | Non-MCE | MCE   | P value | OR    | 95% CI | P value |
|--------------|--------|---------|-------|---------|-------|--------|---------|
| n = 370     | n = 299 | n = 71  |       |         |       |        |         |
| Age, mean (SD) | 67.2 (11.9) | 67.2 (12.1) | 67.2 (11.5) | 0.967 | 0.960 | 0.934-0.987 | 0.004 |
| Male sex, n (%) | 210 (56.8) | 174 (58.2) | 36 (50.7) | 0.252 |       |        |         |
| Medical history, n (%) | | | | | | | |
| Hypertension | 252 (68.1) | 199 (66.6) | 53 (74.6) | 0.188 |       |        |         |
| Diabetes mellitus | 74 (20.0) | 59 (19.7) | 15 (21.1) | 0.792 |       |        |         |
| Atrial fibrillation | 185 (50.0) | 149 (49.8) | 36 (50.7) | 0.895 |       |        |         |
| TOAST, n (%) | | | | | | | |
| LAA | 126 (34.1) | 106 (35.5) | 20 (28.2) | 0.392 |       |        |         |
| CE | 204 (55.1) | 163 (54.5) | 41 (57.7) |       |       |        |         |
| Others | 40 (10.8) | 30 (10.0) | 10 (14.1) |       |       |        |         |
| Baseline data | | | | | | | |
| Baseline SBP, mean (SD) | 142.1 (23.0) | 141.0 (22.5) | 146.5 (24.7) | 0.119 |       |        |         |
| Baseline DBP, mean (SD) | 81.9 (14.2) | 81.3 (13.7) | 84.4 (16.1) | 0.134 |       |        |         |
| Baseline NIHSS, median (IQR) | 17 (13–20) | 16 (13–20) | 18 (16–22) | < 0.001 | 1.076 | 1.016–1.140 | 0.013 |
| Baseline ASPECT, median (IQR) | 9 (8–10) | 9 (8–10) | 8 (8–10) | 0.033 |       |        |         |
| OTP, median (IQR) | 247 (201–300) | 245 (200–300) | 270 (210–310) | 0.328 |       |        |         |
| OTR, median (IQR) | 333 (277–390) | 329 (277–382) | 352 (290–420) | 0.015 |       |        |         |
| Occlusion site, n (%) | | | | | | | |
| ICA | 17 (4.6) | 16 (5.4) | 1 (1.4) | < 0.001 |       |        |         |
| ICA-T | 111 (30.0) | 74 (24.7) | 37 (52.1) |       |       |        |         |
| MCA M1 | 219 (59.2) | 187 (62.5) | 32 (45.1) |       |       |        |         |
| MCA M2 | 23 (6.2) | 22 (7.4) | 1 (1.4) |       |       |        |         |
| Tandem, n (%) | 64 (17.3) | 42 (14.0) | 12 (16.9) | 0.540 |       |        |         |
| Collateral circulation, n (%) | | | | | | | |
| Grade 0 | 71 (19.2) | 41 (13.7) | 30 (42.3) | < 0.001 | 0.360 |        | 0.005 |
| Grade 1 | 163 (44.1) | 132 (44.1) | 31 (43.7) |       | 0.127 |        | < 0.001 |
| Grade 2 | 136 (36.7) | 126 (42.2) | 10 (14.0) |       | 0.051–0.315 |        |         |
| Laboratory test, mean (SD) | | | | | | | |
| FBG, mmol/l | 7.5 (3.3) | 7.1 (3.0) | 9.1 (4.0) | < 0.001 | 1.180 | 1.086–1.281 | < 0.001 |
| LDL, mmol/l | 2.4 (0.9) | 2.4 (0.8) | 2.4 (1.3) | 0.993 |       |        |         |
| BUN, mmol/l | 6.3 (2.5) | 6.2 (2.2) | 6.9 (3.6) | 0.036 |       |        |         |
| Cr, umol/l | 83.2 (40.5) | 81.9 (34.2) | 88.7 (60.6) | 0.208 |       |        |         |
| Treatment, n (%) | | | | | | | |
| Bridging treatment | 86 (23.2) | 65 (21.7) | 21 (29.6) | 0.160 |       |        |         |
| Type of procedure |       |       |       | 0.153 |       |        |         |
| Stent retriever first | 313 (84.6) | 249 (83.3) | 64 (90.2) |       |       |        |         |
| Aspiration first | 18 (4.9) | 14 (4.7) | 4 (5.6) |       |       |        |         |
| Angioplasty or stent first | 39 (10.5) | 36 (12.0) | 3 (4.2) |       |       |        |         |
| Rescue treatment | 67 (18.1) | 47 (15.7) | 20 (28.2) | 0.014 |       |        |         |
| DHC | 18 (4.9) | 0 (0.0) | 18 (25.4) | < 0.001 |       |        |         |
| Outcome, n (%) | | | | | | | |
| mTICI, 2b-3 | 272 (73.5) | 234 (78.3) | 38 (53.5) | < 0.001 | 0.352 | 0.185–0.669 | 0.001 |
| 90-day mRS (0–2) | 167 (45.1) | 162 (54.2) | 5 (7.0) | < 0.001 |       |        |         |
| 90-day mortality | 81 (21.9) | 38 (12.7) | 43 (60.6) | < 0.001 |       |        |         |

Note: MCA occlusion between segments C1–C6.  
Abbreviations: MCE, malignant cerebral edema; TOAST, Trial of Org 10,172 in Acute Stroke Treatment; LAA, large-artery atherosclerosis; CE, cardio-embolism; SBP, systolic blood pressure; DBP, diastolic blood pressure; NIHSS, National Institutes of Health Stroke Scale; ASPECT, Alberta Stroke Program Early CT; OTP, symptom onset to groin puncture time; OTR, time from stroke onset to recanalization; ICA, internal carotid artery; MCA, middle cerebral artery; FBG, fast blood glucose; LDL, low-density lipoprotein; BUN, blood urea nitrogen; Cr, creatinine; DHC, decompressive hemicraniectomy; mTICI, modified thrombolysis in cerebral infarction; mRS, modified Rankin Scale.
a total score and converted into a percentage representing an individual risk of the development of MCE after EVT. The details of the new nomogram are shown in Figure 2. For example, a patient with age at 65 years, blood glucose at > 12.0 mmol/L, baseline NIHSS score at 20.0, mTICI 2b-3, and grade 0 of collateral circulation would have a total of 160.0 points (26.0 points for age, 42.0 points for blood glucose, 32.0 points for baseline NIHSS score, 34.0 points for grade 0 collateral circulation and 36.0 points for mTICI 2b-3).

Figure 2: The nomogram model for predicting the probability of malignant cerebral edema in patients after thrombectomy. The variables, including age, baseline NIHSS score, FBG, the grade of collateral circulation and the degree of TICI, could predict the risk of MCE in patients after thrombectomy. The vertical line of the “Points” axis determines the number of points of every variable for predicting the development of MCE. Add the number of points from each variable. Mark this sum on the “Total point” axis, and draw a vertical line down to meet the “The risk of MCE” axis, to find the patient’s probability of developing MCE.

Abbreviations: mTICI, modified Thrombolysis in Cerebral Infarction; MCE, malignant cerebral edema; NIHSS, National Institutes of Health Stroke Scale.

Figure 3: The receiver operating characteristic curve (A) and the calibration plot (B) of the nomogram model. The dashed line is the reference line where an ideal nomogram would lie. The dotted line is the performance of the nomogram, while the solid line corrects for any bias in the nomogram.
0 points for mTICI 2b-3, and 60.0 points for collateral circulation). The predicted MCE was approximately 54.5% for this patient.

The discrimination of the nomogram was measured by calculating the AUC-ROC value, which was 0.805 (95% CI: 0.750–0.860, Figure 3A). A generally accepted approach suggests that an AUC value of more than 0.75 reflects clearly useful discrimination.21 Also, the ROC curve analysis demonstrated the optimal cutoff value of total nomogram score as a MCE indicator was estimated to be 115.0, which yielded a sensitivity of 77.5% and a specificity of 70.2%. The model was internally validated using 1000 bootstrap samples to calculate the discrimination with accuracy, and the good predictive performance of the nomogram was also confirmed, yielding a notable AUC of 0.805. The Hosmer–Lemeshow goodness-of-fit test showed good calibration of the nomogram (P = 0.681). Additionally, the calibration plot for the nomogram model showed adequate agreement between predictors calculated with the nomogram and actual MCE risk (Figure 3B).

Clinical Outcomes of Malignant Cerebral Edema
The results of univariate analysis of the 90-day favorable outcome are reported in Table S2. In the multivariate analysis, the presence of MCE was the major predictor of poor outcome after EVT treatment (OR: 13.489, 95% CI: 3.322–54.771, P < 0.001). We also found that patients with MCE had an increased mRS score (P < 0.001, Figure 4).

Discussion
In the present study, we found that the occurrence of MCE after EVT was an independent risk factor for poor clinical outcome at 90 days.16 Moreover, our study suggested that several clinical factors, including age, baseline NIHSS score, collateral circulation, FBG, and recanalization status after the procedure, were significantly associated with the development of MCE.

Consistent with the results of previous studies, our findings showed that a higher NIHSS score was an independent predictor for MCE after EVT. The severity of the neurological deficit on admission assessed by the NIHSS score has been suggested to be a stronger predictor of stroke outcome. Moreover, previous studies showed that a higher NIHSS score was associated with cerebral edema after stroke.15,17 Additionally, a higher NIHSS score may imply poor collateral status and a larger infarct core, which are important risk factors for MCE.22,23

Recanalization is strongly associated with improved clinical outcomes after EVT. Our study showed that successful recanalization could significantly decrease the development of MCE. Although abrupt reperfusion is associated with secondary brain damage in animal models of ischemic stroke,24,25 recent studies have confirmed that there is no significant lesion growth after successful recanalization in patients treated with EVT.26 Furthermore, cerebral edema reduction could increase the clinical benefit of thrombectomy in AIS patients.11,27

In the present study, we found that FBG was a powerful predictor of MCE after EVT. Several studies have shown that hyperglycemia could affect clinical outcomes after EVT.28,29 However, the exact mechanism is still unclear. One of the possible mechanisms is that hyperglycemia may increase matrix metalloproteinase-9 activity and disrupt the blood-brain barrier, which may also lead to the occurrence of MCE in patients with AIS.30

Notably, we did not find a significant difference in age between the MCE and non-MCE groups in the univariate analysis. However, since a recent meta-analysis showed that age is a reliable predictor of MCE in ischemic stroke,15 we still included age in the regression analysis in the current study. Unexpectedly, we found that old age was a protective factor for MCE after EVT, which was consistent with the results in the previous meta-analysis. We speculated that the possible reason was that age-related brain atrophy may provide buffering space for brain swelling.

An accurate estimate of the development of MCE after EVT may provide important information to clinicians in clinical decisions and prognostic assessments. Hence, in previous studies, several scoring systems were established to predict the development of MCE in patients with AIS, including those treated with IVD.18,31,32 However, there is no suitable scoring system for patients treated with EVT.
To our knowledge, the present study may be the first attempt to develop a nomogram model for predicting the development of MCE after EVT.

Nomograms are important tools for modern medical decision-making. Compared with previous scoring systems, such as the EDEMA score and DASH score,\textsuperscript{18,31} the nomogram model can assign a probability of developing MCE after EVT, which often provides better-individualized assessments that facilitate management-related decisions. Additionally, the nomogram has higher accuracy and better discrimination ability and is more convenient to use. In the present study, the nomogram model incorporated with five easily available variables had a favorable discrimination ability, as supported by the AUC-ROC value of 0.805 and the optimal calibration curves. Thus, the nomogram model can facilitate the early identification of patients who may develop MCE and support rational decision-making for early DHC of patients with a very high probability of MCE after EVT.

In the present study, we first provided a nomogram model to predict MCE after EVT. However, our study has some limitations. First, this nomogram was based on retrospective data from only 2 institutions. Missing data may affect the accuracy of the model. Second, this study only included a subset of patients who were eligible for EVT according to current guidelines, which may lead to selection bias. Also, the model cannot be generalized to other patient cohorts. Third, known neuroradiologic predictors, such as infarct core, were not available in our study.\textsuperscript{23} Future studies are warranted to investigate whether the integration of novel markers would help improve the accuracy of nomogram prediction.

The nomogram composed of age, baseline NIHSS score, collateral circulation, fast blood glucose level, and recanalization status may predict the risk of MCE in ischemic stroke patients after EVT treatment. Future studies are warranted to externally validate our nomogram in different populations.

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Disclosure
The authors have no conflicts of interest.

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