Hospital Volume Threshold Associated with Higher Survival after Endovascular Recanalization Therapy for Acute Ischemic Stroke

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Background and Purpose Endovascular recanalization therapy (ERT) is becoming increasingly important in the management of acute ischemic stroke (AIS). However, the hospital volume threshold for optimal ERT remains unknown. We investigated the relationship between hospital volume of ERT and risk-adjusted patient outcomes.

Methods From the National Health Insurance claims data in Korea, 11,745 patients with AIS who underwent ERT from July 2011 to June 2016 in 111 hospitals were selected. We measured the hospital’s ERT volume and patient outcomes, including the 30-day mortality, readmission, and postprocedural intracranial hemorrhage (ICH) rates. For each outcome measure, we constructed risk-adjusted prediction models incorporating demographic variables, the modified Charlson comorbidity index, and the stroke severity index (SSI), and validated them. Risk-adjusted outcomes of AIS cases were compared across hospital quartiles to confirm the volume-outcome relationship (VOR) in ERT. Spline regression was performed to determine the volume threshold.

Results The mean AIS volume was 14.8 cases per hospital/year and the unadjusted means of mortality, readmission, and ICH rates were 11.6%, 4.6%, and 8.6%, respectively. The VOR was observed in the risk-adjusted 30-day mortality rate across all quartile groups, and in the ICH rate between the first and fourth quartiles (P<0.05). The volume threshold was 24 cases per year.

Conclusions There was an association between hospital volume and outcomes, and the volume threshold in ERT was identified. Policies should be developed to ensure the implementation of the AIS volume threshold for hospitals performing ERT.

Keywords Stroke; Thrombectomy; Risk adjustment; Hospitals, high-volume; Quality and outcomes

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Introduction

Ischemic stroke is one of the leading causes of mortality and morbidity. Endovascular recanalization therapy (ERT) represents a new treatment paradigm for acute ischemic stroke (AIS), and has been examined in seven large-scale studies. The American Heart Association/American Stroke Association guideline for the treatment of AIS was revised in 2018, and ERT is considered as important as administration of intravenous alteplase in the treatment of AIS. The hospital volume of ERT has been increasing in Korea since 2015. Many studies on ERT have examined the patient aspect of treatment, such as decision making (whom and when to treat or not), technical challenges (incapability of access to the occlusive lesion), and the choice of device (aspiration device versus stentriever). However, relatively little research has examined the systemic aspect of providers.

Several studies have shown that hospital volume could be a surrogate marker of outcomes for various surgical conditions and medical procedures. Other studies have demonstrated that volume-outcome relationship (VOR) is dependent on disease entity, the type of procedure, and type of outcome measurement. Some studies have shown that volume can be an important marker of ERT success in AIS. Eligibility for the Thrombectomy-Capable Stroke Centers certification by the Joint Commission requires a hospital volume of more than 15 ERT cases per year. However, to our knowledge, no studies have investigated whether this volume threshold is appropriate. Using the concept of VOR, it is possible to estimate the minimum volume of ERT for successful treatment of AIS.

We hypothesized that there would be a difference in clinical outcomes between low-volume and high-volume centers that implement ERT for patients with AIS. We also tried to estimate the volume threshold and identify the existing modifiable structures and processes in the high- and low-volume hospitals.

Methods

Study framework

This study was divided into three parts. The first part was the preparatory process before using the administrative database (DB); we validated ERT patients using an identification method and validated a risk-adjusted mortality model incorporating important clinical variables for ischemic stroke (Figure 1A and B). The second part was to examine VORs by analyzing the association between hospital ERT volume and risk-adjusted outcomes derived from summing the individual-level expected outcomes using the administrative DB (Figure 1C). The third part was to calculate the hospital volume threshold (Figure 1D).

Three clinical outcome variables, namely 30-day mortality, 30-day readmission, and postprocedural intracranial hemorrhage (ICH), were selected based on literature review. After identifying other variables for building risk adjustment models for three outcomes from the administrative DB for hospital profiling, we requested and purchased these variables and other related claims data from the Health Insurance Review and Assessment Service (HIRA) in Korea. Next, we verified the administrative DB with the real-world data (clinical DB I) from two university hospitals: Dong-A University Hospital (DAUH) and Pusan National University Yangsan Hospital (PNUYH). We also considered proxy values to replace the National Institutes of Health Stroke Scale (NIHSS), which is difficult to obtain directly from the administrative DB. We used the stroke severity index (SSI) (Supplementary Table 1). To achieve this, we obtained the clinical DB II (Figure 1B) and the local administrative claims DB of all ischemic stroke patients from the aforementioned university hospitals. The derivation and validation of the proxy values were performed separately using different DBs.

From the administrative DB in the HIRA (the HIRA DB), risk-adjusted outcome models for each of the three outcomes were validated separately using clinical DBs.
derived by using multiple regression analysis. Many variables, including modified Charlson comorbidity index (mCCI), and SSI were also analyzed in the model derivation (Supplementary Table 2). The mCCI is one of the variables often used for risk adjustment purposes in the studies using administrative DBs. We measured C-statistics to measure the discrimination power for each of the three models. Next, these models were applied to the HIRA DB, and the outcomes were measured. The relationship between the procedural volume per hospital and the outcomes were then plotted and analyzed to identify VOR (Figure 1C). Based on the results of the VOR analysis, the volume threshold was measured. Finally, to investigate the modified structures and processes, a stratified analysis with hospital characteristics was performed (Figure 1D).

All analyses were conducted with the SAS Enterprise Guide statistical software package version 7.13 (SAS Institute Inc., Cary, NC, USA).

Databases and study populations
To gain access to HIRA DB, we submitted an Institutional Review Board approval document from PNUYH to HIRA. There were many datasets including diagnostic codes of cerebrovascular diseases (such as cerebral ischemic stroke and subarachnoid hemorrhage) and various procedural codes (for instance, mechanical and chemical thrombectomy, Levin tube insertion, and coil embolization of intracranial aneurysm.) From these claims data, 12,013 patients with cerebral AIS who underwent ERT from July 2011 to June 2016 in 162 hospitals were selected. Information about patient and hospital were accessed in a deidentified state.

Clinical DB I was a prospective clinical cohort for patients with AIS undergoing ERT in DAUH and PNUYH. The enrollment period was the same as in the HIRA DB. Clinical DB II was a cohort of all patients with AIS in addition to ERT patients during the same enrollment period and linked with the local claims database of the same patients (Figure 1A and B).

Validity of identifying ERT patients using an administrative database
Although the HIRA DB was used in a de-identified state, it did provide geographical information (city and district). Combining this information with the relative volume of ERT, we produced the DAUH and the PNUYH cohorts from the HIRA DB. To validate these cohorts, a 2x2 contingency table was created, and Clinical DB I was defined as the gold standard. Pearson’s chi-square test and Student’s t-test were used to compare the administrative and clinical databases (Methods section in Supplementary appendix).

Validation of stroke severity index
To derive the SSI, the association between NIHSSs of patients with AIS and SSI were analyzed for all patients with AIS of the DAUH (prediction model). Subsequently, the model was validated with the PNUYH data. The correlation coefficients were 0.70 for the prediction model and 0.68 for the validation model (Supplementary Table 3). The correlation coefficients of both models were similar to those in a previous paper and usually considered relatively high. However, “intensive care unit stay” showed no significant statistical power (Supplementary Table 4). Multiple linear regression analysis was used for creating the SSI model, and Pearson’s chi-square test was used for validating the model (Figure 1B, Methods section in Supplementary appendix).

Analysis

Risk-adjusted outcome models
We operationally defined three outcome variables based on previous research. The 30-day mortality was defined as ‘all deaths from any cause in any hospital within 30 days after initial admission and the ERT procedure.’ Patients who were admitted to multiple hospitals during their AIS episode were assigned to the hospital that performed the ERT procedure.

Thirty-day readmission was defined as ‘readmission to the same hospital from any cause within 30 days after initial admission and the ERT procedure.’ Symptomatic ICH was defined as ‘no pre-existing ICH code before the ERT procedure, and an occurrence of ICH within 30 days after the initial admission and ERT procedure.’

To examine the association between hospital ERT volume and clinical outcomes, risk-adjusted outcome prediction models were built using a multi-level analysis method (Methods section in the Supplementary appendix).

The evidence of distinction between individual-level and hospital-level variables was based on the fact that the random effect on the combined hospital-level outcome was different. In the case of NIHSS, for example, if NIHSS values of patients receiving thrombectomy in different hospitals are distributed relatively equal to each other, adjusting this variable will have a relatively small impact on patient outcomes. We set the hospital-level variable to be able to influence the prognosis of the patient, and at the same time, to be able to function differently according to each hospital.

Simple and multiple logistic regression analyses were performed to select appropriate parameters of risk adjustment modeling. To make final risk adjustment modeling, the multi-level logistic linear regression method was used. C-statistics were used for validation of the risk adjustment model.
Volume-outcome relationships analysis
We examined VORs according to the standard hospital profiling approach used by the Centers for Medicare and Medicaid Services (CMS) of the United States. The approach was as follows: (1) the crude rates for each of the three outcomes were obtained on an individual patient basis; (2) each risk adjustment model was created and applied to each individual patient; (3) following the same method, the individual patient’s expected outcome variable (for instance, risk-adjusted 30-day mortality chance, risk-adjusted 30-day readmission chance) was calculated; (4) outcome values were summed on a hospital basis; and (5) these summed hospital-level outcome values were divided by the hospital volumes. Univariate analyses were performed for all individual and hospital-level variables to determine which variables were incorporated in the multi-level model and examine the problem of multicollinearity.

Using the ratio between the crude hospital-level outcome and the expected hospital-level outcome, the observed to expected ratio (O/E ratio) was calculated and multiplied by the total average outcome rate. This result was defined as the final risk-adjusted outcome value of each hospital. We chose the O/E ratio rather than the predicted to expected ratio (P/E ratio) because the assumption of this P/E ratio method is that there is no VOR. Pearson’s and Spearman’s correlation analyses were used for VOR distribution, and Pearson’s chi-square test was used for quartile group analyses (Figure 1C).

Estimation of hospital volume threshold
The hospital volume threshold was calculated by examining the relative effect on the adjusted outcome odds for an increasing hospital volume of 10 patients for a given hospital size, and varying the given hospital size from an annual volume from boundary value between quartile groups, increments of one case per year. Ten cases were selected as a standardized increment. The rationale for the use of 10 cases as a reference is the relative value of the results of Ross et al. The volume threshold was set to the point at which the odds for an outcome approached 1 as the total volume fell. This method was taken from Ross et al., and was slightly modified for this study.

To calculate the volume threshold, we sorted all the hospitals’ 5-year volumes, divided them into four groups, and measured the relationships between volume and outcomes. Before sorting hospitals by their volume, a substantial proportion of hospitals (51 hospitals) were excluded for two reasons, the first reason being no record of cases between July 2015 and June 2016, and the second being a total number of cases that was less than 20 over 5 years. Restricted cubic spline regression was used for volume threshold estimation. Fisher’s exact test was used to analyze the relationship between hospital volume group and the presence of stroke unit and teaching status (Figure 1D).

Results
Baseline characteristics of the administrative database
Baseline characteristics are shown in Table 1. Of the 111 hospitals included in the VOR analysis, the ERT volumes corresponding to half of all patients were performed in 21 hospitals. Age and the proportion of comorbidities appeared to be higher in the first quartile group than in the other groups.

A total of 51 hospitals (31.4%) were excluded in the VOR analysis due to very low hospital volumes. VOR analyses were conducted on the total number of cases for the remaining 111 hospitals, which was 11,745 (97.7% of the original 12,013 cases). The sample was divided into four groups to have roughly the same number of patients in each group. About 2,900 cases from the most voluminous hospitals were categorized as the first quartile (more than 61 cases per year), the second quartile (31 to 60 cases per year), the third quartile (19 to 30 cases per year), and the fourth quartile (less than 18 cases per year) (Table 1).

Hospital volumes analyses
Over 5 years, 12,013 ERT procedures were performed (Table 1). Some patients (0.009%) had undergone the procedure twice (n=111) or three times (n=2). Of the 162 hospitals, 21 did not perform the ERT procedure between July 2015 and June 2016. The mean volume was 14.8 cases per year, and 59 hospitals (38%) had a volume that was higher than the mean. The overall 30-day mortality rate was 11.6%, 30-day readmission rate was 4.6%, and postprocedural ICH rate was 8.6%.

Validating ERT patient identification
The results of validating ERT patient identification were a sensitivity of 83.2%, a specificity of 98.8%, and a positive predictive value of 80.2% (Figure 1A). To evaluate the discrepancies, in-depth analyses were performed for the 115 true positives and 42 false negatives of the PNUYH cases (Supplementary Tables 5 and 6).

Validation of risk-adjusted outcome models
To solve the multicollinearity problem, other variables included in the SSI were excluded from the univariate analysis, and the statistical model was applied. In the case of mortality, the variables applied to the model were age, SSI, mCCI, sex, hypertension, and atrial fibrillation. The C-statistic of the final mortality rate model was 0.8146, which was higher than the initial tar-
get value of 0.8. Similarly, the same modeling was done for the readmission rate, and included variables were SSI, mCCI, age, transient ischemic attack, prior stroke, and female sex. The C-statistic was 0.6052, which was relatively low. For postprocedural ICH rate, individual-level variables, such as age, SSI, intracranial arterial stenosis, myocardial infarction, hypertension, and transient ischemic attack were selected. 'Teaching status: resident' was included at the hospital-level. The final model C-statistic was 0.6841 (Supplementary Figures 1-3).

Volume-outcome relationships analyses

For mortality rate comparison between quartile groups, the odds ratio (OR) showed differences between the 1st and 3rd quartiles, and between the 1st and 4th quartiles in the crude outcome model ($P<0.05$). However, in the adjustment model, the OR showed differences among the groups from the 2nd to the 4th quartile (1st vs. 3rd, $P=0.003$; 1st vs. 4th, $P<0.001$). Similar results were obtained on comparing readmission and postprocedural ICH rates among quartile groups (Table 2).

In the distribution analysis, three outcomes of both crude and risk-adjusted models showed negative correlations, but there were no statistically significant results (Supplementary Figures 4-6).

Estimation of volume threshold and evaluation of modifiable 'structures and processes'

In the between quartile comparison of hospital volume, there was a significant difference in the OR between the 1st and 2nd quartiles for the risk-adjusted mortality rate, and between the 1st and 4th quartile for the risk-adjusted postprocedural ICH rate. Based on this, we concluded that there was a VOR for ERT in patients with AIS, and subsequently estimated the volume threshold.

It was assumed that the threshold would exist between the volume of the boundary point between the 2nd quartile (less than 60 cases per year) and the boundary point of the 4th quartile (more than 19 cases per year). Between 60 and 19 cases per year, we assessed volume threshold using a modified method presented in a previous paper. In this process, a reversed J-shaped curve was created, and the threshold was set to the point at which the odds for mortality approached "1.0" as the total volume falls. Thus, at the highest value of the J-curve, 22 cases per year corresponded to the inflection point, and 24 cases per year corresponded to where the odds fell below 1. Based on this, we set 24 cases per year as the VOR threshold of ERT for AIS (Figure 2).

To identify modifiable 'structures and processes,' additional

| Variable | All patients | 1st Q group | 2nd Q group | 3rd Q group | 4th Q group |
|----------|--------------|-------------|-------------|-------------|-------------|
| No. of hospitals (included in analyses)* | 162 (111) | 7 | 14 | 25 | 116 (66) |
| No. of patients (included in analyses)* | 12,013 (11,777) | 2,970 | 2,942 | 2,927 | 3,175 (2,938) |
| Age (yr) | 71 (61–78) | 72 (64–78) | 71 (61–77) | 70 (60–77) | 70 (60–77) |
| Male sex | 6,735 (56.06) | 1,607 (54.10) | 1,635 (55.57) | 1,684 (57.53) | 1,675 (57.01) |
| Stroke severity index | | | | | |
| Median (IQR) | 10.33 (6.89–13.95) | 10.22 (6.41–13.95) | 10.33 (7.59–13.95) | 10.48 (7.48–13.95) | 12.11 (8.6–15.01) |
| Range | 2.86–18.74 | | | | |
| Prior stroke | 5,78 (46.51) | 1,450 (48.82) | 1,318 (44.79) | 1,284 (43.86) | 1,426 (48.53) |
| Hypertension | 3,692 (31.34) | 983 (33.09) | 875 (29.74) | 840 (28.69) | 994 (33.83) |
| Atrial fibrillation | | | | | |
| Pre-diagnosed | 2,441 (20.72) | 669 (22.52) | 553 (18.79) | 579 (19.78) | 640 (21.78) |
| After 90 days | 7,258 (61.62) | 1,885 (63.46) | 1,745 (59.31) | 1,755 (59.95) | 1,873 (63.75) |
| Diabetes mellitus | 8,003 (67.95) | 2,146 (72.25) | 1,761 (59.85) | 2,024 (69.14) | 2,072 (70.52) |
| Charlson comorbidity index | 2 (1–2) | 2 (1–2) | 2 (1–2) | 2 (1–2) | 2 (1–3) |
| Hyperlipidemia | 9,281 (78.80) | 2,553 (85.95) | 2,253 (76.58) | 2,128 (72.70) | 2,347 (79.88) |
| ICAS | 1,707 (14.49) | 527 (17.74) | 391 (13.29) | 393 (13.42) | 396 (13.47) |
| Myocardial infarction | 6,035 (51.24) | 1,510 (50.84) | 1,410 (47.92) | 1,396 (47.69) | 1,719 (58.50) |

Values are presented as median (interquartile range) or number (%). Q, quartile; IQR, interquartile range; ICAS, intracranial artery stenosis.

*Substantial proportion of hospitals were excluded in volume-outcome relationship analyses due to very low-volume (no cases in 2015 or less than 20 cases per 5 years).
analyses were conducted regarding the two hypothesized hospital-related variables, stroke unit, and teaching status. Of the 111 hospitals included in the analysis, 50 hospitals (45%) had stroke units, and teaching status hospitals accounted for 79% of all hospitals. The proportions of hospitals with stroke units and teaching status decreased from the 1st to 4th quartile group, and was statistically significant in the fourth quartile group. Due to dichotomization of the classification according to the value determined by the volume threshold, significant differences were found in the proportion of both stroke unit and teaching hospitals between the two quartiles (Table 3).

### Table 2. Relationship between the hospital volume quartile and the odds ratio of 30-day mortality, readmission, and intracranial hemorrhage rates of endovascular recanalization therapy in acute ischemic stroke, 2011–2015 (n=111)

| Quartile | cOR  | 95% CI       | P   | aOR  | 95% CI       | P   |
|----------|------|--------------|-----|------|--------------|-----|
| 30-day mortality rate |      |              |     |      |              |     |
| 1st Q    | 1    | -            | -   | 1    | -            | -   |
| 2nd Q    | 1.07 | 0.91–1.27    | 0.385 | 1.18* | 1.00–1.39    | 0.048 |
| 3rd Q    | 1.17* | 1.00–1.38   | 0.047 | 1.27* | 1.08–1.50    | 0.003 |
| 4th Q    | 1.36* | 1.16–1.60   | <0.001 | 1.37* | 1.16–1.61    | <0.001 |
| 30-day readmission rate |      |              |     |      |              |     |
| 1st Q    | 1    | -            | -   | 1    | -            | -   |
| 2nd Q    | 1.07 | 0.83–1.39    | 0.567 | 0.94 | 0.74–1.20    | 0.649 |
| 3rd Q    | 1.38* | 1.08–1.76   | 0.009 | 0.91 | 0.71–1.16    | 0.469 |
| 4th Q    | 1.24* | 0.96–1.59   | 0.089 | 0.94 | 0.74–1.20    | 0.640 |
| Intracranial hemorrhage rate |      |              |     |      |              |     |
| 1st Q    | 1    | -            | -   | 1    | -            | -   |
| 2nd Q    | 0.98 | 0.81–1.19    | 0.892 | 1.08 | 0.90–1.30    | 0.379 |
| 3rd Q    | 1.08 | 0.89–1.29    | 0.458 | 1.12 | 0.93–1.34    | 0.223 |
| 4th Q    | 1.26* | 1.06–1.57   | 0.009 | 1.21* | 1.01–1.45    | 0.034 |

1st quartile: >61 cases/year; 2nd quartile: 31 to 60 cases/year; 3rd quartile: 19 to 30 cases/year; 4th quartile: <18 cases/year.
cOR, crude odds ratio; CI, confidence interval; aOR, adjusted odds ratio; Q, quartile.
*P<0.05.

### Discussion

In this study, we calculated the minimum hospital volume of ERT for AIS required to maintain minimum treatment quality. Volume threshold was calculated using a spline regression method and was identified to be 24 cases per year. By using the threshold value, hospitals could adhere to the healthcare policies concerning ERT. However, finding the volume requirement alone does not reflect quality, but could be a proxy value.

There are two classical theories of why a VOR is present. The first is the idea of "practice-makes-perfect," and the second is "selective-referral pattern." These theories seem to have been relevant in various previous studies. In our study, it is difficult to distinguish which of the two theories is most relevant. Some researchers have suggested that the relationship between volume and "structure and process," and the relationship between outcome and "structure and process" should be further analyzed, which will allow the actual working mechanism underlying VOR to be identified.

In the VOR analyses, we found no relationship between readmission and hospital volume. It can be assumed that a high readmission is indicative of a poor prognosis. However, readmission in the current study may have been due to the need to perform endarterectomy or carotid artery stenting after the acute phase management of stroke. In Korea, if the treatment period is extended, the patient may be discharged from the
hospital and re-hospitalized due to technicalities surrounding the insurance system. Therefore, it is possible that the high re-admission rate was not related to prognosis.

By using the threshold value identified in this study, stroke centers may be able to better adhere to the healthcare policies concerning ERT. Indeed, such thresholds can guide the designation of a comprehensive stroke center by setting up the manpower and facilities required for the investment. For example, additional stroke centers could be placed in an area where an overtasked comprehensive stroke center is already located, which would allow more than 24 ERT cases to be taken on annually. There is a method to encourage programs in high-volume hospitals; for example, stroke units. This is also associated with qualitative improvement in overall stroke management. If patients are concentrated in a high-volume center but can be dispersed to surrounding centers, the overall outcomes in the region are likely to be better.

Conversely, some studies have shown that when patients were transferred to a high-volume center, the results were better than those from a low-volume center. However, we do not agree with this interpretation because if patients are concentrated in one hospital, it is likely that surgery or other procedures are only required in emergency situations in low-volume centers. It is possible that the overall stroke care system’s quality would be lowered in such situations due to increased transfer time of the patient.

A previous study defined the cut-off value of a high-volume center as 50 cases per year. However, there seemed to be no scientific basis for this threshold. Another study investigated the correlation between the ERT procedure volume and the outcome in AIS, and defined 10 procedures as the cut-off value by dividing centers into low and high-volume. However, they found no VOR after risk adjustment. These studies are also outdated, and the knowledge and equipment used in ERT have since changed. Therefore, we suggest that these studies be understood contextually in their relevant time period.

The present study has some limitations. First, there is a data reliability problem of hospitals when using administrative data. This limitation was overcome by comparing administrative and clinical databases. Second, the database did not allow us to identify other factors such as door to puncture time, initial the Alberta Stroke program early CT score (ASPECTS), basal collateral blood flow status of the patient, early neurological deterioration, presence of intravenous alteplase, and transference from primary stroke center. Because of this, the statistical power of the prediction models was limited. Third, because we analyzed the data at the level of the hospital, we did not analyze the outcomes by individual interventionist. Despite these drawbacks, trend analysis using administrative data and comparison between hospitals was considered appropriate because it could be a viable way to treat the current data.

### Conclusions

To conclude, we found an association between hospital volumes and risk-adjusted outcomes of ERT. We defined the hospital volume threshold of ERT procedures as 24 cases per year. The risk adjustment method using the SSI was taken from a study conducted in Taiwan that was customized to the Korean medical environment. The trends of volumes and outcomes were traced and utilized as a quality indicator of ERT procedures. Although the volume threshold of ERT can be used as reference values to guarantee good outcomes, other factors are likely to affect the outcome of ERT in patients with AIS. Therefore, this threshold value should be used with caution.
Supplementary materials

Supplementary materials related to this article can be found online at https://doi.org/10.5853/jos.2019.00955.

Disclosure

The authors have no financial conflicts of interest.

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Supplementary appendix

Definitions

Ischemic stroke
Ischemic stroke was defined by the I63 code using the 10th revision of the International Statistical Classification of Diseases and Related Health Problems (ICD-10) code system, and endovascular recanalization therapy (ERT) was defined by the procedure code M6631 (thrombolytic treatment—cerebral thrombus removal) or M6633 (mechanical thrombolysis) in the Health Insurance Review and Assessment Service (HIRA) Code system. In this process, we wanted to exclude stroke cases that occurred in-hospital or those that occurred peri-procedurally (e.g., acute ischemic stroke [AIS] as a complication after coil embolization of aneurysmal subarachnoid hemorrhage).

Methods

Validity of identifying ERT patients using an administrative database
The numbers of true positive cases (true positive), missing cases (false negative), incorrect cases (false positive), and correctly excluded cases (true negative) were identified. The sensitivity (true positives / [false negatives + true positives]), specificity (true negatives / [false positives + true negatives]), and positive predictive values (true positives / [true positives + false positives]) were calculated.24

Patients from each database were matched using the patient’s sex, age, admission date, and discharge date. If discrepancies occurred, false negative or false positive was assigned, and true negative notation was assigned to patients who were present in the clinical database but had not undergone the ERT procedure. Pearson’s chi-square test and Student’s t-test were used to compare the administrative and clinical databases.

Validation of stroke severity index
Because the National Institutes of Health Stroke Scale (NIHSS), an established severity score for AIS were unavailable for all ERT patients of this study, we decided to use the stroke severity index (SSI), the claim data-based severity score after validation. The medical insurance system in Taiwan is similar to that of Korea’s, they both utilize fee-for-service, compulsory insurance system, and cover almost everyone in the nation.31 Hence, we used the same methodology of the Taiwan study in this study. It is known that SSI could be used as the proxy for NIHSS, as they both carry the same weight.25

Seven parameters (airway suctioning, bacterial sensitivity test, general ward stay, intensive care unit stay, nasogastric tube, osmotherapy, urinary catheterization) were also selected from the results of the Taiwan study. We agree with the methodology of selecting these seven parameters. We thought that the coefficient values of each seven parameters could be customized to Korean HIRA database, and multiple linear regression from the clinical database performed (Supplementary Table 1).

Risk-adjusted outcome models
The models incorporated patient variables such as demographic characteristics (age and sex) and clinical characteristics (risk factors for ischemic stroke, modified Charlson Comorbidity Index, and the SSI, and hospital variables such as hospital teaching status, geographical location [metropolitan or non-metropolitan], availability of a stroke unit, and hospital ownership [public or private]).28-30 The SSI was included in the model to consider severity of stroke as a substitute for the NIHSS (Supplementary Table 2).25,27

The identification of the hospitals in a “de-identified” administrative database was performed by combining the location information and the relative volume of ERT. Except for the geographical location, the other three hospital parameters were acquired from the official Korean Hospital Association website, Korean Stroke Association website, and each hospital’s website.
**Supplementary Table 1.** Seven parameters of stroke severity index and its explanation

| Descriptions                  | Explanation                                                                 | Code system/Codes          |
|-------------------------------|-----------------------------------------------------------------------------|----------------------------|
| Airway suctioning             | Suctioning of patient's airway in case of altered mental state /or undergoing anesthesia when patient had ERT procedure | HIRA Code/M0135, M0137     |
| Bacterial sensitivity test    | Almost every patients who have shot intravenous antibiotics, they had this test | HIRA Code/B40xx, B413xx, B414x |
| General ward stay             | Relatively mild stroke patients who admitted from emergency department usually go to general ward first | HIRA Code/ABxx             |
| ICU stay                      | Severe stroke patients who admitted from emergency department almost always go to ICU first | HIRA Code/AJxxx           |
| Nasogastric tube              | Severe stroke patients cannot eat orally by themselves, and this tube is usually inserted | HIRA Code/Q2621, Q2622     |
| Osmotherapy                   | For severe stroke patient whose brain tissue damaged massively, usually osmotherapy prescribed to prevent brain herniation | Main Component Code/14800xBU, 148010BU, 148011BU |
| Urinary catheterization       | Severe stroke patient cannot walk by themselves and usually they void by urinary catheterization | HIRA Code/M0060           |

"x" means various numbers, from 1 to 9.
ERT, endovascular recanalization therapy; HIRA, Health Insurance Review and Assessment Service; ICU, intensive care unit.

**Supplementary Table 2.** Parameters for risk adjustment: individual and hospital level

| Fields                          | Name (ICD–10 codes)                                                                 |
|---------------------------------|-------------------------------------------------------------------------------------|
| Demographics                    |                                                                                     |
| Age, sex                        |                                                                                     |
| Risk factor and modified Charlon comorbidity index items | AMI or CHD or CAD (I21, I22, I23, I34, I35)                                          |
|                                 | Congestive heart failure (I50)                                                      |
|                                 | Peripheral vascular disease (I73)                                                    |
|                                 | Dementia (F00, F01, F02, F03)                                                       |
|                                 | Chronic obstructive pulmonary disease (J40, J41, J42, J43, J44)                     |
|                                 | Connective tissue disease (L94)                                                     |
|                                 | Peptic ulcer (K25, K26, K27)                                                        |
|                                 | Mild liver disease (K73, K74)                                                       |
|                                 | Diabetes mellitus (E10, E11, E12, E13, E14)                                          |
|                                 | Diabetes with end organ disease (E13, E14)                                           |
|                                 | Renal disease (N17, N18, N19)                                                       |
|                                 | Nonmetastatic solid tumor (C41, C46, C47, D00, D01, D02, D04, D05, D06, D07, D09) |
|                                 | Leukemia (C91, C92, C93, C94, C95)                                                   |
|                                 | Lymphoma and multiple myeloma (C81, C82, C83, C84, C85, C86, C90)                  |
|                                 | Metastatic tumor (C77, C78, C79)                                                     |
|                                 | AIDS (B20, B21, B22, B23, B34, R75)                                                  |
|                                 | Prior stroke (I63)                                                                 |
|                                 | Transient ischemic attack (I64)                                                     |
|                                 | Hypertension (I10, I11, I12, I13, I15)                                               |
|                                 | Atrial fibrillation (I48)                                                           |
|                                 | other cardiac disease: DCMP, valvular heart disease, PFO (I42.0, I05, Q21.10)      |
|                                 | Extracranial arterial stenosis (I65)                                                 |
|                                 | Intracranial arterial stenosis (I66)                                                |
|                                 | Dyslipidemia (E78)                                                                 |
| Severity for ischemic stroke    | Stroke severity index (no codes, modelled from other items)                         |
| Hospital level items            |                                                                                     |
| Teaching hospital or not        |                                                                                     |
| Geographical location: whether in metropolitan city or not |                                                                                   |
| Stroke unit or not              |                                                                                     |
| Public or private               |                                                                                     |

ICD–10, 10th revision of the International Statistical Classification of Diseases and Related Health Problems; AMI, acute myocardial infarction; CHD, coronary heart disease; CAD, coronary arterial disease; AIDS, acquired immune deficiency syndrome; DCMP, dilated cardiomyopathy; PFO, patent foramen ovale.
**Supplementary Table 3.** Regression statistics of stroke severity index model

| Statistic                                    | Value |
|----------------------------------------------|-------|
| Multiple regression correlation coefficient  | 0.70  |
| R square                                     | 0.49  |
| Adjusted R square                            | 0.49  |
| Standard error                               | 4.52  |
| Observations                                 | 3.13  |

**Supplementary Table 4.** Multiple linear regression model for the stroke severity index (regression coefficient was 0.685)

| Variable                        | Coefficient | Standard error | P     |
|---------------------------------|-------------|----------------|-------|
| Airway suctioning               | 3.7345      | 0.353          | <0.001|
| Bacterial sensitivity test      | 1.1990      | 0.289          | <0.001|
| General ward stay               | -2.9088     | 0.224          | <0.001|
| ICU stay                        | -0.1125     | 0.322          | 0.519 |
| Nasogastric intubation          | 3.4404      | 0.174          | <0.001|
| Osmotherapy                     | 1.8908      | 0.262          | <0.001|
| Urinary catheterization         | 2.7366      | 0.305          | <0.001|
| Constant                        | 5.8726      | 0.234          | <0.001|

ICU, intensive care unit.

**Supplementary Table 5.** Characteristics of the patients in clinical databases (n=4,932)

| Characteristic | DAUH (n=3,176) | PNUYH (n=1,756) | P     |
|----------------|----------------|-----------------|-------|
| Age            | 67.2±12.6      | 70.0±12.9       | <0.001|
| Female sex     | 1,243 (39.7)   | 713 (40.5)      | 0.557 |
| NIHSS          | 4 (1–10)       | 4 (1–9)         | 0.181 |
| LOS            | 7 (4–11)       | 9 (6–14)        | <0.001|

Values are presented as mean±standard deviation, number (%), or median (interquartile range).

DAUH, Dong-A University Hospital; PNUYH, Pusan National University Yangsan Hospital; NIHSS, National Institutes of Health Stroke Scale; LOS, length of stay.

**Supplementary Table 6.** Comparison of the clinical and administrative databases (n=4,932)

| Variable          | DAUH | PNUYH | Overall |
|-------------------|------|-------|---------|
| Sensitivity (%)   | 97.2 | 73.2  | 83.2    |
| Specificity (%)   | 98.5 | 99.3  | 98.8    |
| Positive predictive value (%) | 70.5 | 92    | 80.2    |

DAUH, Dong-A University Hospital; PNUYH, Pusan National University Yangsan Hospital.
Supplementary Figure 1. (A) Selected risk adjustment variables of 30-day mortality models with each odds ratios and 95% confidence intervals and (B) receiver operating characteristic (ROC) curve. SSI, stroke severity index; CCI, Charlson comorbidity index; HTN, hypertension; ICAS, intracranial arterial stenosis; AF, atrial fibrillation.

Supplementary Figure 2. (A) Selected risk adjustment variables of 30-day readmission models with each odds ratios and 95% confidence intervals and (B) receiver operating characteristic (ROC) curve. SSI, stroke severity index; CCI, Charlson comorbidity index; TIA, transient ischemic attack.
**Supplementary Figure 3.** (A) Selected risk adjustment variables of symptomatic intracerebral hemorrhage models with each odds ratios and 95% confidence intervals and (B) receiver operating characteristic (ROC) curve. SSI, stroke severity index; ICAS, intracranial arterial stenosis; MI, myocardial infarction; HTN, hypertension; TIA, transient ischemic attack.

**Supplementary Figure 4.** Plotted graphs of hospital crude (A) and risk-adjusted (B) 30-day mortality rate and endovascular recanalization therapy volumes for 5 years (n=111). Pearson correlation coefficient of –0.21 (P=0.03). However, in the risk-adjusted 30-day mortality model, the correlation coefficient was –0.03 (P=0.71).
Supplementary Figure 5. Plotted graphs of hospital crude (A) and risk-adjusted (B) 30-day readmission rate and endovascular recanalization therapy volumes for 5 years (n=111). Both crude (−0.13) and risk-adjusted (−0.15) models showed a negative correlation. Both models showed statistically insignificant values (P=0.20 and P=0.14).

Supplementary Figure 6. Plotted graphs of hospital crude (A) and risk adjusted (B) symptomatic intracranial hemorrhage rate and endovascular recanalization therapy volumes for 5 years (n=111). Both crude and risk-adjusted models showed a negative correlation (−0.16 and 0.03) and were statistically insignificant (P=0.11 and P=0.73).