The A²DS² Score as a Predictor of Pneumonia and In-Hospital Death after Acute Ischemic Stroke in Chinese Populations

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Abstract

Background and Purpose
Stroke-associated pneumonia (SAP) is a common complication and an important cause of death during hospitalization. The A²DS² (Age, Atrial fibrillation, Dysphagia, Sex, Stroke Severity) score was developed from the Berlin Stroke Registry and showed good predictive value for predicting SAP. We sought to identify the association between the A²DS² score and SAP, and, furthermore, to identify whether the A²DS² score was a predictor for in-hospital death after acute ischemic stroke in a Chinese population.

Methods
This was a retrospective study. 1239 acute ischemic stroke patients were classified to low A²DS² group (0–4) and high A²DS² score (5–10) group. Primary outcome was in-hospital SAP. Logistic regression analyses were performed to identify the association between the A²DS² score and SAP, and also the association between the A²DS² score and in-hospital death.

Results
The overall incidence rates of SAP and in-hospital mortality after acute ischemic stroke were 7.3% and 2.4%, respectively. The incidence rate of SAP in low and high A²DS² score groups was separately 3.3% and 24.7% (P<0.001). During hospitalization, 1.2% patients in low score group and 7.8% patients in high score group died (P<0.001). Multivariate regression demonstrated that patients in high score group had a higher risk of SAP (OR = 8.888, 95%CI: 5.552–14.229) and mortality (OR = 7.833, 95%CI: 3.580–17.137) than patients in low score group.

Conclusions
The A²DS² score was a strong predictor for SAP and in-hospital death of Chinese acute ischemic stroke patients. The A²DS² score might be a useful tool for the identification of patients with a high risk of SAP and death during hospitalization.
Introduction

Stroke-associated pneumonia (SAP) is a common medical complication after stroke, with rates reported between 5.6% and 37.98% [1–11]. SAP frequently occurs in the first week after stroke onset, especially the first 3 days [1, 3]. Evidence shows that SAP is an important risk factor for mortality after stroke [12–16]; 19.1–26% of post-stroke patients with pneumonia, but only 3.5–5% of patients without pneumonia [5,14,15] die during hospitalization. Moreover, SAP increases length of stay (LOS) and hospitalization costs [5,7,9,15], which increase burdens on family and society.

Risk factors for SAP include older age [17–23], male [19,24], atrial fibrillation [19–23], stroke severity [17–20,22,23], dysphagia [2,17–20,22] and total anterior circulation infarct (TACI) or posterior circulation infarct (POCI) stroke subtypes [20,22]. To effectively evaluate the risk of SAP, several scales have been developed, including Kwon’s pneumonia score [17], Sellars’s predictive model [2], Chumbler’s 3-level scoring system [18], Ji’s AIS-APS (acute ischemic stroke-associated pneumonia score) [20], and Smith’s ISAN score [23]. However, these scoring systems are not widely used in routine clinical practice due to small sample size [17], lack of validation [17,18], retrospective nature [18], the unavailability or delayed availability for obtaining predictors (e.g., low abbreviated mental test scores, chronic obstructive pulmonary disease, congestive heart failure) [2,20] and the inability to incorporate important risk factors (e.g., dysphagia) [21,23].

The A²DS² (Age, Atrial fibrillation, Dysphagia, Sex, Stroke Severity) score is a simple scoring system developed from routinely collected data that was available immediately after hospital admission. It was developed from the Berlin Stroke Registry (BSR) cohort [19] and was subsequently validated using German [19], China [22] and United Kingdom [23] stroke registry data. The previous study used the A²DS² score as a continuous variable to predict SAP [22]. However, in the clinical practice, the dichotomized cutoff point was more convenient. Several studies suggested the A²DS² score’s cutoff was 5 point, which represented maximum Youden index [19,20,22]. The objective of our study is to identify the association between the A²DS² score and SAP, also the association between the A²DS² score and in-hospital death in a Chinese acute ischemic stroke population when the A²DS² score was dichotomized into low (0–4) and high (5–10) score groups.

Patients and Methods

Study Population

Subjects were acute ischemic stroke (AIS) patients admitted to the department of neurology in Guangdong Provincial Hospital of Chinese Medicine between August 2005 and July 2008. Inclusion criteria were: (1) ischemic stroke verified by Computerized Tomography (CT) or Magnetic Resonance Imaging (MRI); and (2) time from symptom onset within 7 days. Patients were excluded if any of the components of the A²DS² score were not available.

Ethics Statement

Our study was approved by the ethic committee (2008 GL-37) of 2nd Affiliated Hospital of Guangzhou University of Chinese Medicine. Our study was a retrospective study and all information was pulled out through the electronic system. There was no consent of patients but all the data was analyzed anonymously.
Data Definitions and Data Extraction
Patients’ records with cerebral infarction or ischemic stroke were obtained from the medical record system. The investigators verified the diagnosis according to the CT/MRI results and excluded patients who had a stroke more than 7 days. Finally, patients who met criteria were enrolled in our study. The following information of the patients were pulled out from the system: (1) demographics (i.e., age and sex); stroke risk factors (i.e., hypertension, diabetes mellitus, dyslipidemia, and atrial fibrillation, including a history of atrial fibrillation or documentation of atrial fibrillation at admission, coronary heart disease (CHD), stroke and TIA history, smoking, and drinking); (2) stroke severity at admission as assessed by the National Institute of Health Stroke Scale score (NIHSS); dysphagia (Kubota’s water swallow test Grade III or higher)[25]; (3) outcomes including pneumonia and death during hospitalization.

The A²DS² score was calculated (Age≥75 years = 1, Atrial fibrillation = 1, Dysphagia = 2, male Sex = 1, stroke Severity, National Institutes of Health Stroke Scale 0–4 = 0, 5–15 = 3, ≥16 = 5)[19] and was dichotomized into low (0–4) and high (5–10) score groups[19,20,22].

Primary and Secondary Outcomes
The primary outcome of our study is SAP and the secondary outcome is death during hospitalization. In this study, SAP was diagnosed based on the original medical documents and re-evaluated by the treating physician according to Mann’s diagnostic criteria[26] for pneumonia based on the presence of ≥3 of the following variables: fever (>38°C), productive cough with purulent sputum, abnormal respiratory examination (tachypnea [>22/min], tachycardia, inspiratory crackles, bronchial breathing), abnormal chest radiograph, arterial hypoxemia (PO₂<70mmHg), and isolation of a relevant pathogen (positive gram stain and culture). Patients who had pneumonia before their stroke were not included. Pneumonia that occurred within the first 72 hours after the stroke onset was defined as early-onset pneumonia (EOP)[1]. The routine procedure of nursing care of stroke patients in our stroke unit included dysphagia screening, oral cavity cleaning, head-up position, and regular chest care, which were all done persistently to prevent SAP. Prophylactic antibiotics were not used to prevent pneumonia.

Statistical Analysis
The data were presented as the mean ±standard deviation (SD), the median with interquartile ranges (IQR), or frequencies with percentages, as appropriate. Continuous variables were analyzed with Student’s t- or Kruskal-Wallis tests and categorical variables were analyzed with Chi-square tests. The A²DS² score was analyzed as both continuous and binary variable. To estimate crude and adjusted odds ratios of the A²DS² score for SAP and in-hospital death, univariate and multivariate logistic regression analyses were performed with SAP and in-hospital death as binary outcome. P values < 0.05 were considered to be statistically significant. In logistic regression analysis, accurate estimation of the discriminant function parameters demands sample size of minimum 20 cases for each predictor variable. All analyses were performed using Empower(R) (www.empowerstats.com, X&Y Solutions, Inc., Boston, MA) and R (http://www.R-project.org).

Results
Clinical Characteristics
Records of 1504 patients with the diagnosis of “ischemic stroke” or “cerebral infarction” were extracted from the electronic medical system. 265 patients were excluded because time from
stroke onset was more than 7 days. At last, 1239 patients were analyzed, among which 90 patients (7.3%) had SAP and 30 patients (2.4%) died during hospitalization (Fig 1).

The patients’ mean age was 69.056±11.662, and 732 patients (59.1%) were men.131 patients (10.6%) had atrial fibrillation, 204 patients (16.5%) had dysphagia symptoms, and the NIHSS median score was 3 (IQR 2–6). The median A²DS² score was 2(IQR1-4).1008 patients (81.4%) were in the low A²DS² score group and 231(18.6%) were in the high A²DS² score groups (Table 1).

The overall incidence rate of SAP and in-hospital mortality was 7.3% and 2.4%, respectively. The proportion of SAP varied from 0.6% to 41.9% in patients with different A²DS² score points. The incidence rate of SAP during the hospitalization in the low and high score groups were 3.3% and 24.7% (P<0.001), respectively. During hospitalization, 1.2% patients in the low and 7.8% patients in the high score group died (P<0.001).

Table 1. Characteristics of the study population (n = 1239).

| Characteristic                        | Total     | Total SAP (n = 90) | Without SAP (n = 1149) | P value |
|---------------------------------------|-----------|-------------------|------------------------|---------|
| Age, Mean (SD)                        | 69.056(11.662) | 68.574(11.773)     | 75.211(7.915)          | <0.001  |
| Age group, ≥75, n (%)                 | 451(36.4)  | 47(52.2)           | 404(35.2)              | 0.001   |
| Sex (Male), n (%)                     | 732(59.1)  | 57(63.3)           | 675(58.7)              | 0.394   |
| Vascular risk factor, n (%)           |           |                   |                        |         |
| Hypertension                          | 797(64.3)  | 61(67.8)           | 736(64.1)              | 0.478   |
| Diabetes mellitus                     | 233(18.8)  | 22(24.4)           | 211(18.4)              | 0.165   |
| Dyslipidemia                          | 67(5.5)    | 4(6.0)             | 63(5.5)                | 0.680   |
| Atrial fibrillation                   | 131(10.6)  | 19(21.1)           | 112(9.7)               | 0.001   |
| Coronary heart disease                | 146(11.8)  | 22(24.4)           | 124(10.8)              | <0.001  |
| Previous stroke /TIA                  | 334(27.0)  | 32(35.6)           | 302(26.3)              | 0.056   |
| Smoking                               | 399(32.2)  | 32(35.6)           | 367(31.9)              | 0.480   |
| Drinking                              | 231(18.6)  | 20(22.2)           | 211(18.4)              | 0.365   |
| Dysphagia, n (%)                      | 204(16.5)  | 63(70.0)           | 141(12.3)              | <0.001  |
| Admission NIHSS score, Median (IQR)   | 3(2–6)     | 6(3–10)            | 3(2–6)                 | <0.001  |
| A²DS² score, Median (IQR)             | 2(1–4)     | 6(3–6)             | 2(1–4)                 | <0.001  |
| A²DS² score group, n (%)              |           |                   |                        | <0.001  |
| low score group (0–4)                 | 1008(81.4)| 33(36.7)           | 975(84.9)              |         |
| high score group (5–10)               | 231(18.6)  | 57(63.3)           | 174(15.1)              |         |

TIA indicates transient ischemic attack; NIHSS, National Institutes of Health Stroke Scale; IQR, interquartile range.
Risk Factors for SAP

The univariate analysis showed that age, atrial fibrillation, dysphagia, admission NIHSS score, history of CHD, and high A²DS² score were risk factors for SAP (Table 2). The further multivariate analysis proved that higher A²DS² score was associated with higher risk of SAP (OR = 1.759; 95%CI, 1.560–1.984) even after adjustment for traditional stroke risk factors. Subjects in high A²DS² score group had higher risk for SAP (OR = 8.888; 95%CI, 5.552–14.229) compared to low A²DS² score group.

Risk Factors for In-hospital Mortality

The univariate analysis showed that hypertension, atrial fibrillation, dysphagia, NIHSS at admission and the A²DS² score were risk factors for death during hospitalization (P<0.05) (Table 3). Multivariate logistic regression showed that the A²DS² score’s OR for in-hospital

### Table 2. Univariate analyses of factors related to SAP.

| Characteristic                  | SAP, n (%) | OR    | 95%CI            | P Value |
|--------------------------------|------------|-------|-----------------|---------|
| Age, ≥75                       | 47(10.4)   | 2.016 | 1.310–3.101     | 0.001   |
| Sex (Male)                     | 57(7.8)    | 1.213 | 0.778–1.892     | 0.395   |
| Atrial fibrillation            | 19(14.5)   | 2.478 | 1.440–4.262     | 0.001   |
| NIHSS at admission             | 90(7.3)    | 1.221 | 1.155–1.290     | <0.001  |
| Dysphagia                      | 63(30.9)   | 16.681| 10.280–27.067   | <0.001  |
| Hypertension                   | 71(7.7)    | 1.180 | 0.746–1.866     | 0.478   |
| Diabetes mellitus              | 22(9.4)    | 1.438 | 0.869–2.379     | 0.157   |
| Dyslipidemia                   | 4(6.0)     | 0.804 | 0.286–2.623     | 0.680   |
| Coronary heart disease         | 22(15.1)   | 2.674 | 1.597–4.479     | <0.001  |
| Previous stroke /TIA           | 32(9.6)    | 1.547 | 0.986–2.430     | 0.058   |
| Smoking                        | 32(8.0)    | 1.176 | 0.750–1.842     | 0.480   |
| Drinking                       | 20(8.7)    | 1.270 | 0.756–2.134     | 0.366   |
| High A²DS² score group         | 57(24.7)   | 9.68  | 6.12–15.30      | <0.001  |

SAP indicates stroke-associated pneumonia; TIA, transient ischemic attack; NIHSS, National Institutes of Health Stroke Scale.

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### Table 3. Univariate analyses of factors related to in-hospital death.

| Characteristic                  | Death, n (%) | OR    | 95%CI            | P     |
|--------------------------------|--------------|-------|-----------------|-------|
| Age, ≥75                       | 12(2.7)      | 1.169 | 0.558–2.450     | 0.679 |
| Sex (Male)                     | 19(2.6)      | 1.202 | 0.567–2.547     | 0.632 |
| Atrial fibrillation            | 10(7.6)      | 4.496 | 2.057–9.827     | <0.001|
| Dysphagia                      | 22(10.8)     | 15.518| 6.805–35.389    | <0.001|
| NIHSS at admission             | 30(2.4)      | 1.179 | 1.079–1.288     | <0.001|
| Hypertension                   | 25(3.1)      | 2.830 | 1.076–7.446     | 0.035 |
| Diabetes mellitus              | 6(2.6)       | 1.081 | 0.437–2.677     | 0.865 |
| Dyslipidemia                   | 4(6.0)       | 2.774 | 0.939–8.192     | 0.065 |
| Coronary heart disease         | 7(4.8)       | 2.343 | 0.987–5.560     | 0.054 |
| Previous stroke/TIA            | 8(2.4)       | 0.985 | 0.434–2.234     | 0.971 |
| Smoking                        | 9(2.3)       | 0.900 | 0.408–1.983     | 0.794 |
| Drinking                       | 6(2.6)       | 1.093 | 0.442–2.706     | 0.847 |
| High A²DS² score group         | 18(7.8)      | 7.014 | 3.329–14.779    | <0.001|

TIA indicates transient ischemic attack; NIHSS, National Institutes of Health Stroke Scale.

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mortality was 1.753 (95%CI, 1.444–2.129) (P<0.001) after adjustment for vascular risk factors. Patients in the high score group had a higher risk of in-hospital death than patients in the low score group (adjusted OR = 7.833; 95% CI, 3.580–17.137) (P<0.001) (Table 4).

**Discussion**

SAP is a common medical complication and an important risk factor for mortality after stroke, with rates reported between 5.6% and 37.98%[1–11]. In our study, the incidence rate of SAP 7.3% was lower than some other studies[20,22], which may be due to two reasons. Firstly, the NIHSS score median in our study was 3 (IQR 2–6), while Li’s [22] and Ji’s studies[20] identified median scores of 4 (IQR 2–6) and 5 (IQR 2–9). Secondly, dysphagia screening was routinely performed among stroke patients in our department which was proved to be effective in prevention of SAP[27]. The present study found that 39 (43.3%) SAP was early-onset pneumonia, which is in line with other studies [1,3], indicating that early-onset pneumonia developed in 43%-79% of SAP patients. This indicates that it is important to identify patients with a high risk of SAP based on routinely collected data immediately upon admission.

The results of our study showed that high A2DS2 score group (5–10) had higher risk of SAP than low A2DS2 score group. According to the results of our study, stroke patients whose A2DS2 score was high should get more attention on SAP and might get early prevention or treatment than patients with low A2DS2 score. Given that dysphagia is one of the most important risk factor for SAP[17–20,22], we are currently running a prospective study for the prevention of SAP through dynamic dysphagia assessment in the AIS patients with a high risk of SAP.

Our study also revealed that the A2DS2 score was an important predictor of in-hospital death. Prior study proved clinicians with expertise in stroke performed poorly compared to a validated tool in predicting the outcomes of patients with an acute ischemic stroke. Use of a prognostic tool may be superior for decision-making following an acute ischemic stroke[28]. Several prognostic scores (e.g. Smith’s risk score[29], Saposnik’s iscore [30] and Myint’s SOAR score[31]) had been developed to predict mortality in Acute Stroke. However, Smith’s risk score and Saposnik’s iscore both consisted of 12 components and were very complex; the positive predictive values of the SOAR score were relatively low. Given that most relevant clinical decisions are usually made in the first few days after admission, it’s important to predict in-hospital death as soon as possible after hospital admission. The A2DS2 score was a simple and convenient scoring system and our study indicated that the A2DS2 score was an important predictor of in-hospital death. Patients in the high score group had a higher risk of in-hospital death compared with patients in the low score group (OR = 7.833; 95% CI, 3.580–17.137).

This study has several strengths. First, our findings demonstrate that the A2DS2 score is a strong predictor for SAP and in-hospital death when the A2DS2 score was dichotomized into

| Table 4. The predictive value of the A2DS2 Score for SAP and in-hospital mortality after AIS. |
|----------------------------------------|----------------|----------------|----------------|----------------|
| Variable | Unadjusted OR (95%CI) | Adjusted OR (95%CI) | Unadjusted OR (95%CI) | Adjusted OR (95%CI) |
| the A2DS2 score | 1.779* (1.583–2.000) | 1.759* (1.560–1.984) | 1.681* (1.401–2.018) | 1.753* (1.444–2.129) |
| the A2DS2 score group | | | | |
| Low score group | Ref | Ref | Ref | Ref |
| High score group | 9.679* (6.122–15.302) | 8.888* (5.552–14.229) | 7.014* (3.329–14.779) | 7.833* (3.580–17.137) |

SAP indicates stroke-associated pneumonia. AIS, acute ischemic stroke.

* P<0.001

a Multivariable logistic regression adjusted for history of dyslipidemia, hypertension, diabetes, stroke or transient ischemic attack, coronary heart disease, smoke and drink.

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low and high score groups. In this case, it is easy and efficient to use the A2DS2 score in clinical practice or clinical trial. Second, to our knowledge, this is the first study to use the A2DS2 score as a predictor of in-hospital death for AIS patients and demonstrates that the A2DS2 score is an important predictor of in-hospital death. Third, though the Centers for Disease Control and Prevention criteria for the diagnosis of pneumonia were not applied, pneumonia in our study is comparable to that of other large studies[19,20], making large quantities of misdiagnoses of SAP unlikely. Moreover, the present study documents the exact date of SAP onset, which demonstrates that the incidence of early-onset pneumonia was high and it’s important to predict SAP as soon as possible after admission.

Our study also had some limitations. First, our study is a retrospective study from a single center, which could have impacted selection bias. Second, because this is an observational study, we cannot rule out the possibility that the results might be affected by some unmeasured confounders such as the use of angiotensin-converting enzyme inhibitors or angiotensin receptor blockers[32], β-Blocker[33] and metoclopramide[34]. Third, in our study, we did not analyze the association between mechanical ventilation and outcome. Ventilator-associated pneumonia is assumed to be caused by mechanical ventilation itself and other risk factors that substantially differ from those identified for the development of SAP[35]. Also, in the process of developing the A2DS2 score, restricting the analysis to patients not ventilated did not change any of the observed association substantially [19].

This study was a retrospective study, thus, there is a need for further prospective validation. We believe that the A2DS2 score is a useful tool for predicting SAP and death during hospitalization, which will aid clinical decision-making for patients who have AIS.

Conclusions
The A2DS2 score was a strong predictor for SAP and in-hospital death in Chinese AIS patients. The A2DS2 score might be a useful tool for identifying patients with a high risk of SAP and death during hospitalization.

Supporting Information
S1 Text. PLOS One Clinical Studies Checklist. (DOCX)
S2 Text. STROBE checklist v4 combined PlosMedicine. (DOCX)

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Author Contributions
Conceived and designed the experiments: YC MZ. Performed the experiments: XZ SY. Analyzed the data: MZ XZ. Contributed reagents/materials/analysis tools: LW RY ML XL GL. Wrote the paper: XZ.

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