Long-Term Mortality Among ICU Patients With Stroke Compared With Other Critically Ill Patients

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Objectives: Assessment of all-cause mortality of intracerebral hemorrhage and ischemic stroke patients admitted to the ICU and comparison to the mortality of other critically ill ICU patients classified into six other diagnostic subgroups and the general Dutch population.

Design: Observational cohort study.

Setting: All ICUs participating in the Dutch National Intensive Care Evaluation database.

Patients: All adult patients admitted to these ICUs between 2010 and 2015; patients were followed until February 2017.

Interventions: None.

Measurements and Main Results: Of all 370,386 included ICU patients, 7,046 (1.9%) were stroke patients, 4,072 with ischemic stroke, and 2,974 with intracerebral hemorrhage. Short-term mortality in ICU-admitted stroke patients was high with 30 days mortality of 31% in ischemic stroke and 42% in intracerebral hemorrhage. In the longer term, the survival curve gradient among ischemic stroke and intracerebral hemorrhage patients stabilized. The gradual alteration of mortality risk after ICU admission was assessed using left-truncation with increasing minimum survival period. ICU-admitted stroke patients who survive the first 30 days after suffering from a stroke had a favorable subsequent survival compared with other diseases necessitating ICU admission such as patients admitted due to sepsis or severe community-acquired pneumonia. After having survived the first 3 months after ICU admission, multivariable Cox regression analyses showed that case-mix adjusted hazard ratios during the follow-up period of up to 3 years were lower in ischemic stroke compared with sepsis (adjusted hazard ratio, 1.21; 95% CI, 1.06–1.36) and severe community-acquired pneumonia (adjusted hazard ratio, 1.57; 95% CI, 1.39–1.77) and in intracerebral hemorrhage patients compared with these groups (adjusted hazard ratio, 1.14; 95% CI, 0.98–1.33 and adjusted hazard ratio, 1.49; 95% CI, 1.28–1.73).

Conclusions: Stroke patients who need intensive care treatment have a high short-term mortality risk, but this alters favorably with increasing duration of survival time after ICU admission in patients with both ischemic stroke and intracerebral hemorrhage, especially compared with other populations of critically ill patients such as sepsis or severe community-acquired pneumonia patients. (Crit Care Med 2020; 48:e876–e883)

Key Words: brain ischemia; critical care; intensive care unit; intracranial hemorrhages; mortality; stroke
Stroke is a major global healthcare issue. Despite the declining occurrence rate of stroke and subsequent mortality over time (1), stroke is still the third leading cause of death in the Netherlands (2) and fifth in the United States (3). Furthermore, stroke is a leading cause of functional disability (4).

Stroke-associated cerebral damage can concomitantly cause compromise of other vital organ functions and, consequently, patients may need treatment in the ICU.

With expanding number of treatment and support options, acute stroke care and the need for intensive care will be increasingly intertwined, and an increasing number of stroke patients will be admitted to the ICU in the future (5, 6).

The broad definition of stroke includes ischemic stroke (IS) and intracerebral and subarachnoid hemorrhage (SAH) (7). In practice, SAH differs from acute IS and intracerebral hemorrhage (ICH) with respect to demographics, recuperation, and effect on other organ function due to catecholamine release (8, 9). Therefore, this research focuses on patients with IS or ICH admitted to the ICU. Hereafter, we will refer to “stroke” as composite outcome for IS and ICH.

Mortality in patients with IS and ICH admitted to the ICU has been investigated before (10–15), but these studies did not include long-term mortality, were limited to small study populations, and did not compare stroke patients to ICU patients admitted for other diagnoses. Earlier research using the Dutch National Intensive Care Evaluation (NICE) database (16) compared large groups of critically ill patients and assessed case-mix adjusted mortality beyond hospitalization in several diagnostic subgroups (17). However, this study did not include patients with IS and ICH. For other diagnostic subgroups, it is recognized that the sequelae of ICU admission extend beyond hospitalization, but this is still unknown for the subset of stroke patients who need intensive care admission, particularly in the recent era in which treatment modalities for (mainly ischemic) stroke have changed markedly (18–24).

In this contemporary era of innovative treatment modalities for stroke, the aim of this study was to assess all-cause mortality from ICU admission onwards of patients with IS and ICH. Furthermore, we compared this to the mortality of the general population and six predefined diagnostic subgroups (including SAH) of ICU patients and investigated the mortality risk patterns over time after having survived the initial event that led to ICU admission.

MATERIALS AND METHODS

Data Sources

In this cohort study, consecutive adult patients admitted to all ICUs participating in the Dutch NICE database between 2010 and 2015, comprising over 90% (85 ICUs in 2015) of ICUs, were considered eligible and recruited (16). Patients were followed using the national medical insurance claim database Vektis (Vektis Beheer BV, Zeist, The Netherlands). The duration of follow-up ended at death or the last date of observation, as documented in the Vektis database until February 2017. Linkage between the NICE and Vektis database was obtained to define date of death after ICU admission using a deterministic linkage algorithm (25). Healthcare insurance is compulsory for all Dutch citizens; hence, the Vektis database includes nearly complete coverage of all medical care in The Netherlands. Exclusion criteria were linkage discrepancies between the NICE and Vektis database and not fulfilling the Acute Physiology and Chronic Health Evaluation (APACHE) IV criteria (26).

The mortality rate for stroke patients in this study was compared with the mortality rate of the general Dutch population (GDP). The GDP mortality rate was generated from StatLine (CBS, The Hague, The Netherlands) (2). In order to standardize indirectly, the GDP weighted average death rates were calculated using age and gender distribution with the selected population of stroke patients as standard. Stroke patients were treated according to the most recent national protocol “Ischemic Stroke and Intracranial Hemorrhage” (27) of the Dutch Society for Neurology with involvement of the Dutch Society of Critical Care.

The NICE database is registered according to the Dutch Data Protection Act. The medical ethics committee of the Amsterdam University Medical Center waived informed consent for this study under Dutch national law (Institutional Review Board protocol number W18_049#18.067).

Diagnostic ICU Subgroups

Stroke patients were defined as patients admitted to ICU due to IS or ICH, not including patients with SAH or deep cerebral venous thrombosis. Other patients were classified into predefined diagnostic subgroups according to the APACHE IV classification system: 1) SAH, 2) traumatic brain injury (TBI), 3) sepsis, 4) severe community-acquired pneumonia (sCAP), 5) cardiac surgery, 6) nonsurgical cancer, and 7) other diagnoses. Detailed definitions are shown in Figure 1. These groups were chosen in order to compare our stroke study population to populations with other neurologic reasons for ICU admissions (1 and 2), reasons for frequent admissions (5), and with known chronic morbidity and sequelae beyond critical care (3, 4, and 6). All ICU patients except stroke patients were analyzed as a combined group as well.

Statistical Analyses

The primary outcome was defined as all-cause short-term (30 d) and long-term (1 yr) mortality after ICU admission.

First, crude cumulative mortality risks were assessed by Kaplan-Meier survival estimates for patients with (I) IS and (II) ICH, (III) all ICU patients except stroke as a combined group, (IV) the GDP and the above predefined diagnostic ICU subgroups (V-1 to V-6).

Second, to assess the gradual alteration of mortality risk with increasing survival periods and to understand the risk pattern after ICU admission, mortality risk was assessed for patients who survived ICU discharge, 30 days, 3 months, and 1 year. This was done by using left-truncation for these predefined time points. Survival of patients who survived up to these time points was analyzed by repeating the Kaplan-Meier
estimates from these points onwards and starting the curve again at 100%.

Third, the differences in mortality between stroke patients and other diagnostic ICU subgroups were analyzed in a multivariable Cox regression model. Case-mix adjusted hazard ratios (HRadj) and corresponding 95% CIs were calculated. The population of IS and ICH were used as reference groups with the same left-truncations as described above. The HRs were adjusted for age, sex, APACHE III score, and calendar year of ICU admission (2010–2015). Correlation between duration of survival of patients admitted to the same ICU was taken into account by including hospital as a random intercept in the models. Preparation of the data files was done using SPSS, Version 24.0 (IBM Corp., Armonk, NY). The statistical analyses were performed using statistical environment R, Version 3.4.3 (R Foundation for Statistical Computing, Vienna, Austria).

RESULTS
During the study period, 483,419 patients were admitted to the participating ICUs. After exclusion of patients not fulfilling the APACHE IV criteria and linkage discrepancies, 370,386 patients were included for analyses. Of the included patients, 7,046 (1.9%) were stroke patients, 4,072 with IS, and 2,974 with ICH (Fig. 1). Baseline characteristics and follow-up data of patients with IS, ICH, all subgroups, and all ICU patients (including stroke) are shown in Table 1. Supplemental Table 1 (Supplemental Digital Content 1, http://links.lww.com/CCM/F616) contains baseline characteristics of additional groups, including patients after cardiac surgery, with nonsurgical cancer and the combined group (all ICU patients except stroke).

Demographics were mainly comparable between IS and ICH. Patients with ICH had higher ICU re-admission rate and lower Glasgow Coma Scale at admission, while patients with IS had more chronic diagnoses. In 30.9% of patients with ICH, a neurosurgical intervention was performed. In total, 3,366 stroke patients died (48%) during the study period with a median (interquartile range) follow-up of 9.6 months (0.2–23.7 mo).
Survival Distribution Compared With GDP and Diagnostic ICU Subgroups

The survival of patients with IS was significantly better compared with patients with ICH (log-rank test $p < 0.001$). In both IS and ICH, crude cumulative survival at the end of follow-up was worse compared with the GDP and the combined group of all other ICU patients (Fig. 2). The inset of Figure 2 shows a large decline in survival of stroke patients within the first 10 days after ICU admission with mortality rate of 29%. This is in contrast to the longer term, in which the survival curve gradient among both IS and ICH patients stabilizes.

Survival in cardiac surgery, TBI and SAH patients, remained better than in stroke during the whole follow-up period (Supplemental Fig. 1, Supplemental Digital Content 2, http://links.lww.com/CCM/F617). The crude cumulative survival of sepsis and sCAP, initially better than IS, became worse compared with IS after approximately 18 months. Among all diagnostic subgroups of our ICU populations, patients with ICH and nonsurgical cancer had the lowest cumulative survival with 1-year survival of respectively 49% and 30% (Supplemental Fig. 1a, Supplemental Digital Content 2, http://links.lww.com/CCM/F617).

Mortality Risk Patterns Over Time Compared With Diagnostic ICU Subgroups

With an increasing minimum of survival time, using left-truncation, differences in survival curves of patients with IS and ICH

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**TABLE 1. Demographics, Length of Stay and Mortality, and Follow-Up in Ischemic Stroke and Intracerebral Hemorrhage Patients and Four of the Six Investigated Other Diagnostic Subgroups (See Supplemental Table 1 [Supplemental Digital Content 1, http://links.lww.com/CCM/F616] for Baseline Characteristics of All Groups)**

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Ischemic Stroke</th>
<th>Intracerebral Hemorrhage</th>
<th>Subarachnoid Hemorrhage</th>
<th>Traumatic Brain Injury</th>
<th>Sepsis</th>
<th>Severe Community-Acquired Pneumonia</th>
<th>All Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>n (%)</td>
<td>4,072 (1.1)</td>
<td>2,974 (0.8)</td>
<td>2,650 (0.7)</td>
<td>4,356 (1.2)</td>
<td>24,189 (6.5)</td>
<td>22,833 (6.2)</td>
<td>370,386</td>
</tr>
<tr>
<td>Age, mean (sd)</td>
<td>67.5 (13.7)</td>
<td>60.7 (15.1)</td>
<td>58.5 (12.9)</td>
<td>53.3 (20.4)</td>
<td>66.9 (14.2)</td>
<td>66.9 (14.5)</td>
<td>63.9 (15.3)</td>
</tr>
<tr>
<td>Sex, men, %</td>
<td>56.5</td>
<td>56.8</td>
<td>35.0</td>
<td>69.9</td>
<td>56.1</td>
<td>59.2</td>
<td>59.5</td>
</tr>
<tr>
<td>Daytime admissions (8:00–18:00 hr, %)</td>
<td>37.5</td>
<td>33.3</td>
<td>35.3</td>
<td>28.6</td>
<td>35.3</td>
<td>36.5</td>
<td>54.9</td>
</tr>
<tr>
<td>One or more chronic diagnoses (%)a</td>
<td>18.1</td>
<td>14.6</td>
<td>9.5</td>
<td>8.4</td>
<td>44.3</td>
<td>59.4</td>
<td>33.5</td>
</tr>
<tr>
<td>Acute Physiology and Chronic Health Evaluation IV predicted mortality, %, mean (sd)</td>
<td>40 (20)</td>
<td>50 (20)</td>
<td>40 (30)</td>
<td>20 (20)</td>
<td>50 (20)</td>
<td>40 (20)</td>
<td>20 (20)</td>
</tr>
<tr>
<td>ICU readmission rate (%)</td>
<td>3.8</td>
<td>6.5</td>
<td>7.6</td>
<td>4.2</td>
<td>6.3</td>
<td>3.3</td>
<td>5.6</td>
</tr>
<tr>
<td>Glasgow Coma Scale at admission, mean (sd)</td>
<td>11.1 (4.4)</td>
<td>8.6 (4.7)</td>
<td>10.7 (4.9)</td>
<td>10.0 (4.7)</td>
<td>13.8 (2.6)</td>
<td>137 (29)</td>
<td>137 (32)</td>
</tr>
<tr>
<td>ICU LOS, d, mean (sd)</td>
<td>2.9 (6.1)</td>
<td>4.6 (8.4)</td>
<td>5.3 (8.0)</td>
<td>5.3 (8.5)</td>
<td>6.2</td>
<td>(10.1)</td>
<td>66 (9.4)</td>
</tr>
<tr>
<td>ICU mortality (%)</td>
<td>696 (17)</td>
<td>907 (31)</td>
<td>580 (22)</td>
<td>566 (13)</td>
<td>4,500 (19)</td>
<td>3,347 (15)</td>
<td>29,305 (8)</td>
</tr>
<tr>
<td>Of which within 7 d (% of total ICU mortality)</td>
<td>605 (86.9)</td>
<td>808 (89.1)</td>
<td>499 (86.0)</td>
<td>414 (73.1)</td>
<td>3,394 (75.4)</td>
<td>2,096 (62.6)</td>
<td>22,515 (76.8)</td>
</tr>
<tr>
<td>Hospital LOS, d, mean (sd)</td>
<td>11.0 (13.2)</td>
<td>13.9 (18.6)</td>
<td>14.3 (15.3)</td>
<td>14.5 (16.6)</td>
<td>207</td>
<td>(22.1)</td>
<td>173 (23.1)</td>
</tr>
<tr>
<td>Hospital mortality, n (%)</td>
<td>1,172 (29)</td>
<td>1,210 (41)</td>
<td>722 (27)</td>
<td>747 (17)</td>
<td>6,236 (26)</td>
<td>5,193 (23)</td>
<td>44,557 (12)</td>
</tr>
<tr>
<td>30-d mortality, n (%)</td>
<td>1,280 (31)</td>
<td>1,263 (42)</td>
<td>748 (28)</td>
<td>776 (18)</td>
<td>6,358 (26)</td>
<td>5,516 (24)</td>
<td>47,796 (13)</td>
</tr>
<tr>
<td>3-mo mortality, n (%)</td>
<td>1,476 (36)</td>
<td>1,401 (47)</td>
<td>817 (31)</td>
<td>881 (20)</td>
<td>7,694 (32)</td>
<td>7,026 (31)</td>
<td>60,220 (16)</td>
</tr>
<tr>
<td>1-yr mortality, n (%)</td>
<td>1,663 (41)</td>
<td>1,519 (51)</td>
<td>856 (32)</td>
<td>987 (23)</td>
<td>9,394 (39)</td>
<td>8,909 (39)</td>
<td>81,396 (22)</td>
</tr>
<tr>
<td>Total amount of deceased patients, n (%)</td>
<td>1,783 (44)</td>
<td>1,583 (53)</td>
<td>879 (33)</td>
<td>1,058 (24)</td>
<td>10,638 (44)</td>
<td>10,480 (46)</td>
<td>99,485 (27)</td>
</tr>
<tr>
<td>Follow-up, mo, mean (sd)</td>
<td>13.3 (12.2)</td>
<td>11.1 (12.2)</td>
<td>14.7 (12.3)</td>
<td>17.5 (11.7)</td>
<td>13.9</td>
<td>(12.0)</td>
<td>13.8 (11.8)</td>
</tr>
<tr>
<td>1-yr follow-up, n (%)</td>
<td>2,084 (51.2)</td>
<td>1,249 (42.0)</td>
<td>1,474 (55.6)</td>
<td>2,936 (67.4)</td>
<td>12,684 (52.4)</td>
<td>11,859 (51.9)</td>
<td>245,350 (66.2)</td>
</tr>
<tr>
<td>2-yr follow-up, n (%)</td>
<td>1,029 (25.3)</td>
<td>661 (22.2)</td>
<td>803 (30.3)</td>
<td>1,616 (37.1)</td>
<td>6,465 (26.7)</td>
<td>5,706 (25.0)</td>
<td>128,568 (34.7)</td>
</tr>
</tbody>
</table>
The mortality risk in ICH was higher compared with IS without left-truncation (HR_{adj} was significantly higher than 1.00) but this difference disappeared with increasing minimum of survival time (i.e., with applying left-truncation the HR_{adj} became close to and nonsignificantly different from 1.00).

The mortality risk was lower in TBI and cardiac surgery and higher in nonsurgical cancer when compared with either IS or ICH, regardless of whether survival periods were increased, so no alteration of mortality risk occurred compared with IS or ICH.

Among SAH patients, the risk for mortality was lower compared with the mortality risk of patients with ICH, regardless of left-truncation. Compared with IS patients, the hazard was not different in SAH patients when no left-truncation was applied (HR_{adj}, 0.98; 95% CI, 0.90–1.07), but became significantly lower in SAH with increasing minimum of survival time.

Among patients with sCAP, the mortality risk after surviving 3 months was significantly higher compared with either IS (HR_{adj}, 1.57; 95% CI, 1.39–1.77) or ICH patients (HR_{adj}, 1.49; 95% CI, 1.28–1.73) (Supplemental Fig. 2, a and b, Supplemental Digital Content 3, http://links.lww.com/CCM/F618).

**DISCUSSION**

This national multicenter study has unraveled the survival pattern of patients with IS and ICH admitted to the ICU in a cohort of over 7,000 stroke patients with a follow-up of up to 3 years. We found that stroke patients admitted to the ICU had high short-term mortality, more pronounced in ICH than in IS when compared with other critically ill patients. However,
when a stroke patient survived the first 30 days after ICU admission (69% in IS and 58% in ICH), the prognosis gradually improved. Furthermore, it was shown that mortality risk after surviving 3 months was approaching statistical significance in patients with ICH and significantly lower in IS, compared with both sepsis and sCAP patients. After having survived 1 year, mortality risk in both ICH and IS were significantly lower compared with sepsis and sCAP.

The 1-year mortality rate of 41% in IS and 51% in ICH patients admitted to the ICU found in this study is comparable to earlier studies (10–15, 28–30), although previous literature did not report mortality beyond one year after ICU admission in such a large cohort. Furthermore, the present study did not only compare stroke patients to other neurologic ICU patients (12–14) but also included cardiac surgery patients with low predicted mortality and patients with diagnoses suspected for long-term sequelae and high predicted mortality, such as ICU patients with sepsis, sCAP, and nonsurgical cancer (31). This has led to the observation that long-term mortality in critically ill stroke patients is higher than in TBI or SAH, as shown earlier (12–14) and that the survival patterns of patients with sepsis and sCAP is totally different from stroke patients.

An explanation for this finding may be that stroke can be seen as a one-time event and is mostly failure of a single organ, whereas sepsis is a lengthier and ongoing syndrome with a wide spectrum of underlying chronic conditions and often accompanied with multiple organ involvement. The proportion of patients with one or more chronic diagnosis in all separate diagnostic subgroups in our study (18% in IS and 14% in ICH compared with 44% in sepsis) underlines this explanation. In addition, recent literature has described that patients surviving sepsis acquire new physical disability and cognitive impairment leading to further health deterioration after hospital discharge with an increased hazard of death and therefore implying persistent long-term mortality risk (32–35).

Stroke leads to evident physical and cognitive disability. However, the data in this study show that in ICU-admitted stroke patients it is important to recognize the concept that patients who survive the first 30 days after suffering from a stroke, have a relative favorable subsequent survival compared with other diseases necessitating ICU admission such as patients admitted due to sepsis or sCAP.

The strengths of this study are its large number of admissions from 85 ICUs across the country, the long-term follow-up, and usage of the existing APACHE IV model, which has been validated in the Dutch ICU population (36). Furthermore, we were able to correct for various known determinants in order to confirm our results on mortality risk in stroke patients compared with the other predefined diagnostic ICU subgroups, since risk of mortality is highly correlated to the underlying case-mix of patients (37).

However, some limitations need to be addressed. This was a cohort study with all-cause mortality as outcome variable and therefore not taking reason of death into account, including withdrawal of care and comfort of care transitions. Therefore, the high all-cause mortality in stroke patients might be related to withdrawal of care in severe cases and selection of patients with less deficits, contributing to the gradual decrease of mortality over time if a patient survived the first period. On the other hand, withdrawal of care does imply clinical deterioration as judged by the members of the healthcare team, so these patients were probably severe stroke patients with an expected high mortality.

Second, our results are limited to the subset of acute stroke patients admitted to the ICU. No extrapolation is possible to IS or ICH patients admitted from the emergency department to the ward or stroke unit due to lack of expected benefit (as judged by the admitting physician) from ICU care on one hand or lack of necessity of ICU care on the other hand. To put our study numbers in perspective, in 2010–2012, approximately 35,000 patients were admitted annually to one of all Dutch hospitals due to suspicion of stroke (20.7–21.1 clinical admissions per 10,000 citizens) (2). More recent, 33,733 stroke patients were registered in 2017 in 71 hospitals within the country. Of patients with IS, 21% received IV thrombolysis and 4.5% underwent intra-arterial thrombectomy (39).

Third, the administrative NICE database creates the possibility to analyze case-mix adjusted long-term mortality far beyond hospitalization in a large and extensive cohort of ICU-admitted stroke patients, but unfortunately does not contain data concerning functional status or posthospital location and therefore this study pertains only to mortality risks of critically ill stroke patients and not to their life’s quality or abilities in daily life. Unfortunately, we could not collect data concerning National Institutes of Health Stroke Scale scores (40, 41) or ICH scores (42) via the NICE database. Furthermore, due to linkage discrepancies, 10.4% of the NICE database records were excluded from the study. However, the deterministic link-age algorithm that was used leads to a small number of false-positive links (43), indicating that our linked data set is reliable.

Finally, since we did not possess data concerning vascular territories of stroke or numbers on reperfusion rates or anticoagulant therapies in respectively IS and ICH, heterogeneity among the study population might have occurred. The unraveled survival pattern in IS and ICH patients admitted to the ICU could be due to homogenous populations with high mortality hazards in short-term and lower in longer term. Another explanation might be that our study group consists of several subgroups with different stroke subtypes within the stroke population with different prognoses and survival patterns.

CONCLUSIONS

Stroke patients who need intensive care treatment have a high short-term mortality risk. However, the mortality risk alters favorably with increasing survival period after ICU admission in patients with both IS and ICH, especially compared with other populations of critically ill patients such as sepsis or sCAP patients. Future research is needed to reveal clinical variables which can identify stroke patients at ICU admission who will survive that first critical period in the ICU. These data can subsequently assist in personalizing critical care and informing patients and their family caregivers about patterns
and probabilities concerning survival and quality of life after ICU admission due to an IS or ICH.

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This work was performed at Amsterdam University Medical Center, Amsterdam, The Netherlands; Leiden University Medical Center, Leiden, The Netherlands; and University Medical Center Utrecht, Utrecht, The Netherlands.

Bart F. Geerts and M. Sesmu Arbous contributed equally to this article.

Mariëlle K. van Valburg, Bart F. Geerts, and M. Sesmu Arbous designed the study protocol. Mariëlle K. van Valburg, Wilson F. Abdo, Walter M. van den Bergh, Janneke Hof, Walther N. K. A. van Mook, Bob Siegerink, Arjen J. C. Slooter, Marieke J. H. Wermers, Bart F. Geerts, and M. Sesmu Arbous created the study project. Fabian Termorshuizen and Sylvia Brinkman analyzed the data and performed the statistical analyses. Mariëlle K. van Valburg and Fabian Termorshuizen wrote the draft, and all coauthors critically revised the article and approved the final version for publication.

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The Study on Outcome and Prognosis in Hemorrhagic and Ischemic Stroke patients admitted To the Intensive Care (SOPHISTIC) project, and the resulting research has received funding from HandicapNL (former Revalidatiefonds) under project number R2015057 for financing the linkage of the National Intensive Care Evaluation database to the Vektis national insurance claim database.

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The National Intensive Care Evaluation database is registered according to the Dutch Data Protection Act. The medical ethics committee of the Amsterdam University Medical Center waived informed consent for this study under Dutch national law (Institutional Review Board protocol number W18.049#18.067).

The Study on Outcome and Prognosis in Hemorrhagic and Ischemic Stroke patients admitted To the Intensive Care (SOPHISTIC) project was registered in the Dutch Trial Registry (https://www.trialregister.nl) as NTR7438.

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REFERENCES


