RESEARCH ARTICLE

Dysphagia in Acute Stroke: Incidence, Burden and Impact on Clinical Outcome

Marcel Arnold1*, Kai Liesirova1*, Anne Broeg-Morvay1, Julia Meisterernst1, Markus Schlager1, Marie-Luise Mono1, Marwan El-Koussy2, Georg Kägi3, Simon Jung1†, Hakan Sarikaya1‡

1 Department of Neurology, University Hospital of Berne, Berne, Switzerland, 2 Department of Neuroradiology, University Hospital of Berne, Berne, Switzerland, 3 Department of Neurology and Neurophysiology, Kantonsspital St. Gallen, St. Gallen, Switzerland

☯ These authors contributed equally to this work.
‡ MA and KL are shared first authors on this work. SJ and HS also contributed equally to this work and are shared last authors on this work.
★ marcel.arnold@insel.ch

Abstract

Background

Reported frequency of post-stroke dysphagia in the literature is highly variable. In view of progress in stroke management, we aimed to assess the current burden of dysphagia in acute ischemic stroke.

Methods

We studied 570 consecutive patients treated in a tertiary stroke center. Dysphagia was evaluated by using the Gugging Swallowing Screen (GUSS). We investigated the relationship of dysphagia with pneumonia, length of hospital stay and discharge destination and compared rates of favourable clinical outcome and mortality at 3 months between dysphagic patients and those without dysphagia.

Results

Dysphagia was diagnosed in 118 of 570 (20.7%) patients and persisted in 60 (50.9%) at hospital discharge. Thirty-six (30.5%) patients needed nasogastric tube because of severe dysphagia. Stroke severity rather than infarct location was associated with dysphagia. Dysphagic patients suffered more frequently from pneumonia (23.1% vs. 1.1%, p < 0.001), stayed longer at monitored stroke unit beds (4.4 ± 2.8 vs. 2.7 ± 2.4 days; p < 0.001) and were less often discharged to home (19.5% vs. 63.7%, p = 0.001) as compared to those without dysphagia. At 3 months, dysphagic patients less often had a favourable outcome (35.7% vs. 69.7%, p < 0.001), less often lived at home (38.8% vs. 76.5%, p < 0.001), and more often had died (13.6% vs. 1.6%; p < 0.001). Multivariate analyses identified dysphagia to be an independent predictor of discharge destination and institutionalization at 3 months, while severe dysphagia requiring tube placement was strongly associated with mortality.
Conclusion

Dysphagia still affects a substantial portion of stroke patients and may have a large impact on clinical outcome, mortality and institutionalization.

Introduction

Dysphagia is a common complication of stroke, but estimates of its frequency vary considerably [1,2]. It is an important cause of pneumonia within the first days after stroke and previous studies reported an increased risk of mortality in the acute phase [3–5]. Furthermore, dysphagia has been shown to be associated with malnutrition, dehydration and increased length of hospital stay [6]. Over the last decades, acute stroke management has substantially improved in many countries through reperfusion techniques, introduction of stroke units and stroke prevention. Accordingly, several studies have reported a substantial decline in both stroke incidence and associated mortality in developed countries [7–9]. Considering these advances in stroke management, we were interested in the current incidence and burden of dysphagia in a comprehensive stroke center and its association with pneumonia, discharge destination and clinical outcome at 3-month follow-up.

Methods

The Inselspital is a tertiary stroke center and the main teaching hospital of the University of Bern, Switzerland, with a catchment area of approximately 2 million inhabitants. The Bernese Stroke Registry is a computer-based system and was set up in January 2000 to prospectively collect data from consecutive acute stroke patients at Inselspital. The study was approved by the Local Ethics Committee Bern. Patients with acute ischemic stroke admitted between January 2012 and November 2013 were retrospectively evaluated. The authors had access to identifying patient information. Data was anonymized prior to author access.

Demographic variables, vascular risk factors, infarct location and the use of thrombolysis were systematically recorded. Stroke territory was divided into anterior and posterior, based on previously used definitions [10,11]. Among strokes in posterior cerebral circulation, brainstem involvement was registered separately as it is supposed to be strongly associated with dysphagia. Stroke severity at admission was assessed with the National Institutes of Health Stroke Scale (NIHSS) score [12]. All patients treated with thrombolysis were admitted to a stroke unit for at least 24 hours. Clinical outcome at 3 months was evaluated by a certified stroke nurse or neurologist using the modified Rankin Scale (mRS) score [13]. The following parameters were also assessed: mortality during hospitalization and at 3 months, discharge destination and living situation at 3 months, length of hospital stay for different units (stroke unit vs. ward), incidences of dysphagia and pneumonia as well as the frequency of antibiotic treatment and nasogastric tube insertion. Pneumonia was diagnosed by the treating physician according to Centers for Disease Control and Prevention (CDC) criteria and based on typical findings in pulmonary examination (tachypnoea, inspiratory crackles, bronchial breathing) or chest radiograph (infiltration, consolidation), and additional evidence of either fever (body temperature >38°C), cough with purulent sputum or pathogen isolation in either blood or sputum culture [14]. Experienced physiotherapists with special training in dysphagia regularly checked the swallowing ability in each stroke patient within 24 hours after admission by using the Gugging Swallowing Screen (GUSS) and oral feeding was withheld until intact swallowing was demonstrated [15]. Dysphagia evaluation started with part 1 of GUSS, which provides one point for...
each of the following 5 criteria: (I) patient is vigilant, (II) normal voluntary coughing, (III) nor-
mal deglutition of saliva (IV) without drooling and (V) without voice change. If patients had a
GUSS score <5 points in part 1, further dysphagia evaluation was performed by additional
investigations (Videofluoroscopic Evaluation of Swallowing (VFES) or Fiberoptic Endoscopic
Evaluation of Swallowing (FEES) or placement of a nasogastric tube, depending on the judg-
ment of the treating therapist and physician. If the maximum of 5 points was achieved in part 1
of GUSS, the next subset (direct swallowing test, part 2) of GUSS was continued consisting of 3
subtests (semisolid swallowing, liquid swallowing, solid swallowing). The highest possible score
in the GUSS is 20 points (5 points in part 1 and 15 points in part 2), denoting normal swallow-
ing ability. For this study, dysphagia was defined as GUSS score of 19 points or less. A score of
less than 10 points was defined as severe dysphagia with high aspiration risk, thus these
patients received a nasogastric tube (“nil by mouth”) [15]. In patients with GUSS score between
10 and 19 points, stepwise special diet was ordered depending on severity of dysphagia [16].

The primary aim of this study was to assess the incidence of dysphagia in stroke patients
and to compare the clinical outcomes at 3 months, namely death and favourable outcome
(defined as mRS 0–1). In addition, we aimed to evaluate predictors of dysphagia and potential
association with following secondary endpoints: occurrence of pneumonia, use of antibiotics
and chest radiographs, length of hospital stay in various units, discharge destination and insti-
tutionalization at 3 months.

**Statistical Analysis**

Normally distributed data were expressed as mean±standard deviation (SD) and compared
using Students t-test. The 2 groups (patients with dysphagia vs. patients with normal swallow-
ing ability) were compared using Mann Whitney U-test for continuous variables and the Pear-
son’s chi-square test for binary variables. Multivariate logistic regression analysis was
performed to assess the independent effects of dysphagia and the other predictors on the out-
come parameters favourable outcome, institutionalization and mortality at 3 months. In a first
step, the influence of every single potential predictor on the outcomes was evaluated using uni-
ivariate logistic regression analysis. The parameters examined were age, baseline NIHSS score,
sex, infarct location (anterior vs. posterior cerebral circulation), brainstem infarction, arterial
hypertension, hypercholesterolemia, diabetes and smoking. In a second step, a multivariate
logistic regression analysis was performed, including all potential predictors with a p-value
<0.2 from univariate analyses. Significance was set at p<0.05 level.

**Results**

A total of 570 consecutive patients with ischemic stroke were included in this study. Mean age
was 65.1 years (range, 19.6–94.7 years), 366 (64.2%) patients were male and thrombolysis was
performed in 156 (27.3%) patients. Ischemic stroke affected the anterior cerebral circulation in
436 (76.5%) patients, posterior cerebral circulation in 104 (18.3%) and both territories in 28
(4.9%). Exact stroke localization could not be established in 2 patients. Brainstem infarction
was present in 82 of 104 (78.9%) patients with posterior circulation stroke.

Dysphagia was diagnosed in 118 (20.7%) patients (mean age, 65.6; range, 23.5–91.0 years).
The clinical characteristics of patients with and without dysphagia are shown in Table 1.
Patients with dysphagia had more severe neurological deficits at baseline (mean NIHSS score
9.7±7.0 vs. 4.5±5.1; p<0.001) and had less often hypertension as compared to patients without
dysphagia (36.2% vs. 63.7%; p = 0.029). Other clinical characteristics were similar in both
groups.
Baseline characteristics of dysphagic patients without tube placement and those with tube insertion are compared in Table 2. Patients who required tube feeding had higher NIHSS scores (13.4±6.8 vs. 8.2±6.5; p<0.001) and were more often women (48.7% vs. 27.2%; p = 0.022). Age was not significantly different between groups (68.9±15.2 vs. 64.0±14.0; p = 0.091).

There were no major differences in the frequency of dysphagia stratified to stroke location: dysphagia was observed in 95 (21.8%) patients with anterior circulation stroke, 18 (17.3%) with posterior circulation stroke, and 5 (17.9%) with both anterior and posterior circulation stroke. Of 82 patients with brainstem infarction, 20 (24.4%) had dysphagia. The frequency of dysphagia at hospital discharge was similar in anterior, posterior and brainstem stroke (10.8%, 8.7%, and 13.4%, respectively). Dysphagia persisted in 4 of 5 patients with both anterior and posterior circulation stroke.

Tables 3 and 4 summarize the severity and persistence of dysphagia among stroke patients with swallowing dysfunction. During hospitalization, 36 of 118 (30.5%) patients with dysphagia needed a nasogastric tube and one patient required a percutaneous endoscopic gastrostomy.
At hospital discharge, dysphagia persisted in 60 of 118 (50.9%) patients who had dysphagia at admission. Tube feeding was ongoing in 23 (19.5%) patients at discharge. In-hospital pneumonia occurred in 27 (22.9%) patients with dysphagia and in 5 (1.1%) without dysphagia (p<0.001, see Fig 1). Accordingly, the use of antibiotics was markedly higher in patients with dysphagia than those without dysphagia (28.0% vs. 4.6%, p<0.001). Multivariate logistic regression analysis showed that dysphagia (OR, 27.4; 95% CI, 10.2–73.7; p<0.001) was independently associated with pneumonia. Compared to dysphagic patients without tube insertion, those receiving tube feeding had much higher risk for in-hospital pneumonia, need of antibiotic treatment and death at 3 months (35.1% vs. 17.3%, p = 0.035, 51.4% vs. 17.3%, p<0.001, and 27.0% vs. 7.4%, p = 0.004, respectively). However, after adjustment for confounders, the association between tube placement and pneumonia was no longer statistically significant (OR, 2.2; 95% CI, 0.89–5.5; p = 0.087).

Patients with dysphagia more often underwent chest radiographs (0.5±0.9 vs. 0.2±0.9 per person, p = 0.004) during hospital stay and were admitted to the stroke unit longer (4.4±2.8 vs. 2.7±2.4 days, p<0.001) as compared to patients without dysphagia. The total duration of hospital stay was similar in both groups (7.9±4.8 vs. 7.2±4.4 days, p = 0.145). Distinctive differences were observed with respect to discharge destination (Fig 1): patients with dysphagia had a higher probability for transfer to a rehabilitation clinic and a lower chance to be discharged home as compared to non-dysphagic patients (78.0% vs. 35.4%, p = 0.001 and 19.5% vs. 63.7%, p = 0.001, respectively). At 3 months, patients presenting with dysphagia were still less likely to live at home than patients with normal swallowing (38.8% vs. 76.5%, p<0.001).

In-hospital mortality was slightly higher in dysphagic patients (2.5% vs. 0.7%, p = 0.075), but at 3 months, mortality clearly increased in patients with dysphagia (13.6% vs. 1.6%, p<0.001). The frequency of favourable outcome was lower in dysphagic patients (35.6% vs. 69.7%, p<0.001) and half (50.0% vs. 21.4%, p<0.001) were living at a healthcare unit or rehabilitation clinic (Fig 1).

Multivariate logistic regression analyses identified stroke severity measured by NIHSS to be an independent predictor of dysphagia (odds ratio [OR], 1.1; 95% confidence interval [CI], 1.1–1.2; p<0.001). Discharge destination was significantly affected by dysphagia (OR, 4.7; 95% CI, 2.7–7.9; p<0.001) and stroke severity (OR, 1.1; 95% CI, 1.1–1.2; p<0.001).

Predictors of mortality at 3 months were severe dysphagia requiring tube feeding (OR, 8.5; 95% CI, 2.8–26.2; p<0.001), in-hospital pneumonia (OR, 9.7; 95% CI, 3.2–29.4; p<0.001), and brainstem involvement (OR, 4.1; 95% CI, 1.4–12.3; p = 0.011). Severe dysphagia requiring tube

### Table 3. Severity of dysphagia among 118 patients with swallowing disorder.

|                        | %      | n     |
|------------------------|--------|-------|
| Dysphagia requiring tube feeding | 31.3   | 37/118|
| Dysphagia requiring nasogastric tube feeding | 30.5   | 36/118|
| Dysphagia requiring PEG* | 0.8    | 1/118 |

* PEG stays for percutaneous endoscopic gastrostomy

doi:10.1371/journal.pone.0148424.t003

### Table 4. Persistence of dysphagia among 118 patients with swallowing disorder.

|                        | %      | n     |
|------------------------|--------|-------|
| Persisting dysphagia at hospital discharge | 50.9   | 60/118|
| Persisting dysphagia requiring tube feeding at discharge | 19.5   | 23/118|

doi:10.1371/journal.pone.0148424.t004
feeding, persisting dysphagia at hospital discharge and baseline stroke severity were inversely associated with favourable outcome at 3 months (OR, 0.3; 95% CI, 0.1–0.9; p = 0.028; OR, 0.2; 95% CI, 0.1–0.4; p<0.001; and OR, 0.96; 95% CI, 0.93–0.99; p = 0.016, respectively). The following factors were independently associated with institutionalization at 3 months: dysphagia (OR, 3.1; 95% CI, 1.7–5.5; p<0.001), stroke severity (OR, 1.2; 95% CI, 1.2–1.3; p<0.001) and brainstem involvement (OR, 0.3; 95% CI, 0.1–0.8; p = 0.008).

**Discussion**

In this study, dysphagia affected more than one in five patients with ischemic stroke and was independently associated with in-hospital pneumonia, discharge destination and institutionalization, while severe dysphagia was a strong predictor of unfavourable outcome and mortality at 3 months. Furthermore, dysphagia had a significant impact on increased healthcare consumptions such as chest radiographs or antibiotics.
The reported frequency of dysphagia after stroke varied considerably in earlier studies, ranging from 50% to 80% [1,17–20]. In our study, the incidence of dysphagia was substantially lower than these estimates. These variations can be attributed to differences in definition and assessment of dysphagia, timing of swallowing examination, and patient selection. Advances in acute stroke treatment and management at a stroke unit may also have contributed to the lower rate in this study. Nevertheless, the frequency of swallowing dysfunction in our study still accounts for a substantial proportion, thus a regular and precise screening in patients with acute stroke is advisable.

An important finding of this study is the large impact of dysphagia on clinical outcome at 3 months. Patients with dysphagia had an 8.5-fold higher risk of death as compared to those with normal swallowing. Furthermore, dysphagia was independently associated with institutionalization of stroke patients at 3 months. In addition, multivariate analyses showed tube placement due to severe dysphagia was an independent predictor of mortality and unfavourable outcome. These observations are in line with other studies reporting increased mortality, less favourable outcome and higher institutionalization rates at 3 months in stroke patients presenting with dysphagia [3,4]. Stroke severity was another significant determinant for clinical recovery and residency at 3 months. Of note, mortality at 3 months was associated more strongly with severe dysphagia than with stroke severity at baseline. These data indicate that dysphagia has a large effect on survival, clinical recovery, and dependency after stroke. The use of thrombolysis was higher in patients presenting with dysphagia. This probably relates to the fact that dysphagia is associated with severe strokes and that thrombolysis is used more often in severe cases. Similar findings were reported by Okubo and colleagues [21].

It is well known that stroke is associated with high in-hospital costs and our study indicates that dysphagia may additionally increase the medical expenses. Affected patients more often underwent chest radiographs and antibiotic treatment, stayed longer at a stroke unit and more frequently were transferred to rehabilitation clinics. On the other hand, the overall duration of hospital stay was not significantly different in patients with normal swallowing compared to those with dysphagia, which contradicts results from earlier studies [22–24]. This difference may be explained by our policy to transfer stable stroke patients at an early stage to primary care hospitals in the Bernese stroke network. However, dysphagic patients stayed significantly longer at stroke unit as compared to those without dysphagia. A recent study from South Carolina reported that dysphagia after stroke significantly increases medical costs, driven by higher hospital and durable medical equipment expenses [24]. The 1-year costs of post-stroke dysphagia was estimated to be $4’510, as compared to $1’703 attributable to post-stroke aphasia [24,25].

Patients with dysphagia had a much higher risk for pneumonia as compared to patients with normal swallowing in our study, which may further increase the hospitalization costs [26]. This finding is consistent with a review that indicated a 3-fold increased risk of pneumonia in patients with dysphagia and an 11-fold increased risk in those with aspiration [1]. Furthermore, pneumonia predicted increased risk of death at 3 months. This observation is in line with a large multi-center study reporting an increased 30-day and 1-year mortality due to stroke-associated pneumonia [27]. The incidences of post-stroke pneumonia in intensive care units range from 22% to 47%, depending on study design and definition of chest infection [28,29]. In comparison, the overall rate of pneumonia in our study population was rather low (6%). Regular dysphagia screening in stroke units by trained therapists may explain this, as this strategy has been shown to decrease the incidence of pneumonia in acute stroke patients [30,31]. On the other hand, still one in four stroke patients with dysphagia developed pneumonia despite regular monitoring. This observation calls for further improvement in the management of dysphagic patients. Recent studies indicate that functional magnetic or electrical...
stimulation may improve dysphagia as compared to conventional swallowing therapy, but these results need to be confirmed by larger studies [32,33]. There could be a need for such novel approaches, as swallowing dysfunction may persist in many patients for a longer time [34,35]. In our study, half of all patients with initial dysphagia still had dysphagia at hospital discharge.

Our study indicates that initial stroke severity is the main risk factor for dysphagia as shown by previous studies [3,21]. Stroke location was not associated with dysphagia in our study. This finding contradicts the assumption that dysphagia may be mainly caused by posterior circulation or brainstem strokes. Accordingly, studies have shown that total anterior circulation strokes have the highest frequency of dysphagia and lesions of the frontal and insular cortex independently predict prolonged dysphagia after stroke [3,22,36,37]. Supratentorial strokes such as lesions of pre-motor or motor cortices and basal ganglia may cause dysphagia by affecting the planning and execution of swallowing or pharyngeal peristalsis, whereas brainstem lesions may impair oro-pharyngeal sensation, laryngeal elevation and timing of pharyngeal swallow [2,38]. Rates of recovery from dysphagia were comparable in anterior and posterior circulation stroke in our study. Furthermore, simultaneous lesions in anterior and posterior circulation stroke were associated with an unfavourable prognosis, but the low number of patients prohibits any firm conclusions. Similar to our findings, stroke territory did not influence the prognosis of dysphagia in other studies [17,36]. Poor recovery has been reported in severe strokes and especially large lesions of primary motor cortex, whereas dysphagia after brainstem stroke had a favourable outcome on long-term [20,22,39]. Patients with brainstem infarction were at lower risk for institutionalization in our study. We believe that this may be mainly associated with the severity of stroke, which is supposed to be less severe in brainstem involvement. Accordingly, several studies have shown that patients with posterior circulation stroke have lower National Institutes of Health Stroke Scale (NIHSS) scores than patients with anterior circulation stroke [40–42]. In addition, large infarcts in territory of middle cerebral artery (MCA) often cause severe hemiparesis, cognitive impairment, aphasia or neglect requiring intense rehabilitation and / or institutionalization. In contrast, in brainstem infarctions higher cortical functions are not affected and many patients with brainstem infarct and without pyramidal tract involvement do not have paresis (e.g. Wallenberg’s syndrome). Therefore, patients with brainstem infarctions may be at lower risk for institutionalization at 3 months. Finally, selection bias may be another cause for this observation: patients with severe brainstem strokes generally have markedly higher mortality rates and we may have followed just survivors with small brainstem infarcts at 3 months, whereas hemicraniectomy in severe MCA stroke often decrease mortality without improving clinical recovery (survivor bias).

Another finding of our study is that patients with nasogastric tube placement had a much higher risk of death as compared to dysphagic patients without tube insertion, whereas the association with the occurrence of pneumonia was rather weak in multivariate analyses. Langdon and colleagues reported significantly higher rates of respiratory infection in strictly tube-fed stroke patients as compared to those fed orally [43]. On the one hand, one may argue that tube placement is probably a marker of increased aspiration risk as this was performed in patients with large infarcts and severe dysphagia (mainly GUSS score <10 points). However, it is still under debate whether placement of nasogastric tubes additionally increases the risk of pneumonia by promoting colonization of oropharynx with pathogenic bacteria [44,45], as most aspiration pneumonia are assumed to result from bacterial origin [46]. Noteworthy, feeding tubes do not prevent aspiration of gastroesophageal reflux [44,47]. A retrospective review indicated nasogastric tubes and immobility to be stronger predictors for respiratory infections than dysphagia in acute stroke [48]. These data suggest that the benefit and harm of nasogastric tubes needs to be investigated in further studies. Stringent oral hygiene, attention to upright
positioning during enteral feeding, mobilization of bedridden patients and routine change of nasogastric tubes may decrease the risk of pneumonia [43,49–51].

This study has several limitations. The retrospective study design is the main drawback, although dysphagia, baseline data, risk factors and clinical outcomes were assessed routinely in patients. GUSS has been shown to be a reliable method for dysphagia screening, however, no consensus exists on the optimal diagnostic tool [52]. Another limitation is the use of NIHSS score as the only marker of stroke severity, whereas volume measurements of infarct lesions were not performed due to variations in types of imaging. Furthermore, classification of dysphagia was rather broad as it was dichotomized between dysphagia and severe dysphagia only. In addition, we did not have data on pre-existing comorbidity that may have resulted in swallowing impairment prior to stroke. In view of broad confidence intervals, the results must be interpreted cautiously and need to be confirmed by larger studies. Finally, our findings may not be generalized with respect to our monocentric study design in a tertiary care stroke center.

In summary, this study indicates that dysphagia still affects a substantial portion of stroke patients. Dysphagia was clearly associated with clinical outcome, mortality and healthcare expenses despite advanced stroke treatment. In view of this burden, our data call for further research to improve management of dysphagia in acute stroke.

Author Contributions
Conceived and designed the experiments: MA SJ HS. Analyzed the data: KL SJ MA HS. Wrote the paper: HS MA KL SJ. Made critical revision to the manuscript: AB-M JM MS M-LM ME-K GK. Collected data: KL AB-M JM MS M-LM ME-K GK. Wrote grant for funding: SJ MA HS.

References
1. Martino R, Foley N, Bhogal S, Diamant N, Speechley M, Teasell R. Dysphagia after stroke: incidence, diagnosis, and pulmonary complications. Stroke J Cereb Circ. 2005; 36: 2756–2763. doi:10.1161/01.STR.0000190056.76543.eb
2. González-Fernández M, Ottenstein L, Atanelov L, Christian AB. Dysphagia after Stroke: an Overview. Curr Phys Med Rehabil Rep. 2013; 1: 187–196. doi:10.1007/s40141-013-0017-y PMID: 24977109
3. Paciaroni M, Mazzotta G, Corea F, Caso V, Venti M, Milia P, et al. Dysphagia following Stroke. Eur Neurol. 2004; 51: 162–167. doi:10.1159/000077663 PMID: 15073441
4. Smithard DG, Smeeton NC, Wolfe CDA. Long-term outcome after stroke: does dysphagia matter? Age Ageing. 2007; 36: 90–94. doi:10.1093/ageing/afi149 PMID: 17172601
5. Smithard DG, O'Neil PA, Parks C, Morris J. Complications and outcome after acute stroke. Does dysphagia matter? Stroke J Cereb Circ. 1996; 27: 1200–1204.
6. Altman KW, Yu G-P, Schaefer SD. Consequence of dysphagia in the hospitalized patient: impact on prognosis and hospital resources. Arch Otolaryngol Head Neck Surg. 2010; 136: 784–789. doi:10.1001/archoto.2010.129 PMID: 20713754
7. Carandang R, Seshadri S, Beiser A, Kelly-Hayes M, Kase CS, Kannel WB, et al. Trends in incidence, lifetime risk, severity, and 30-day mortality of stroke over the past 50 years. JAMA. 2006; 296: 2939–2946. doi:10.1001/jama.296.24.2939 PMID: 17190894
8. Lee S, Shafe ACE, Cowie MR. UK stroke incidence, mortality and cardiovascular risk management 1999–2008: time-trend analysis from the General Practice Research Database. BMJ Open. 2011; 1: e000269. doi:10.1136/bmjopen-2011-000269 PMID: 22021893
9. Fang MC, Coca Perraillon M, Ghosh K, Cutler DM, Rosen AB. Trends in stroke rates, risk, and outcomes in the United States, 1988 to 2008. Am J Med. 2014; 127: 608–615. doi:10.1016/j.amjmed.2014.03.017 PMID: 24680794
10. Tatu L, Moulin T, Bogousslavsky J, Duvernoy H. Arterial territories of the human brain: cerebral hemispheres. Neurology. 1998; 50: 1699–1708. PMID: 9633714
11. Bogousslavsky J, Van Melle G, Regli F. The Lausanne Stroke Registry: analysis of 1,000 consecutive patients with first stroke. Stroke J Cereb Circ. 1989; 18: 1083–1092.
12. Brott T, Adams HP, Olinger CP, Marler JP, Barsan WG, Biller J, et al. Measurements of acute cerebral infarction: a clinical examination scale. Stroke J Cereb Circ. 1989; 20: 864–870.
van Swieten JC, Koudstaal PJ, Visser MC, Schouten HJ, van Gijn J. Interobserver agreement for the assessment of handicap in stroke patients. Stroke J Cereb Circ. 1988; 19: 604–607.

14. Garner JS, Jarvis WR, Emori TG, Horan TC, Hughes JM. CDC definitions for nosocomial infections, 1988. Am J Infect Control. 1988; 16: 128–140. PMID: 341893

15. Trapl M, Enderle P, Nowotny M, Teuschl Y, Matz K, Dachenhausen A, et al. Dysphagia bedside screening for acute-stroke patients: the Gugging Swallowing Screen. Stroke J Cereb Circ. 2007; 38: 2948–2952. doi: 10.1161/STROKEAHA.107.483933

16. Crary MA, Mann GDC, Groher ME. Initial psychometric assessment of a functional oral intake scale for dysphagia in stroke patients. Arch Phys Med Rehabil. 2005; 86: 1516–1520. doi: 10.1016/j.apmr.2004.11.049 PMID: 16084801

17. Mann G, Hankey GJ, Cameron D. Swallowing disorders following acute stroke: prevalence and diagnostic accuracy. Cerebrovasc Dis Basel Switz. 2000; 10: 380–386. doi:16094

18. Nilsson H, Ekberg O, Olsson R, Hindfelt B. Dysphagia in stroke: a prospective study of quantitative aspects of swallowing in dysphagic patients. Dysphagia. 1998; 13: 32–38. doi: 10.1007/PL00009547 PMID: 9391228

19. Kidd D, Lawson J, Nesbitt R, MacMahon J. The natural history and clinical consequences of aspiration in acute stroke. QJM Mon J Assoc Physicians. 1995; 88: 409–413.

20. Meng NH, Wang TG, Lien IN. Dysphagia in patients with brainstem stroke: incidence and outcome. Am J Phys Med Rehabil Assoc Acad Physiatr. 2000; 79: 170–175.

21. Okubo PCM, Fábio SRC, Domenis DR, Takayanagui OM. Using the National Institute of Health Stroke Scale to predict dysphagia in acute ischemic stroke. Cerebrovasc Dis Basel Switz. 2012; 33: 501–507. doi:10.1159/000336240

22. Broadley S, Croser D, Cottrell J, Creevy M, Teo E, Yiu D, et al. Predictors of prolonged dysphagia following acute stroke. J Clin Neurosci Off J Neurosurg Soc Australas. 2003; 10: 300–305.

23. Falsetti P, Acciai C, Palilla R, Bosi M, Carpineti F, Zingarelli A, et al. Oropharyngeal dysphagia after stroke: incidence, diagnosis, and clinical predictors in patients admitted to a neurorehabilitation unit. J Stroke Cerebrovasc Dis Off J Nati Stroke Assoc 2009; 18: 329–335. doi: 10.1016/j.jsctd.2009.01.009

24. Bonilha HS, Simpson AN, Ellis C, Mauldin P, Martin-Harris B, Simpson K. The one-year attributable cost of post-stroke dysphagia. Dysphagia. 2014; 29: 545–552. doi: 10.1007/s00545-014-9543-8 PMID: 24948438

25. Ellis C, Simpson AN, Bonilha H, Mauldin PD, Simpson KN. The one-year attributable cost of poststroke aphasias. Stroke J Cereb Circ. 2012; 43: 1429–1431. doi: 10.1161/STROKEAHA.111.647339

26. Wilson RD. Mortality and cost of pneumonia after stroke for different risk groups. J Stroke Cerebrovasc Dis Off J Nati Stroke Assoc. 2012; 21: 61–67. doi: 10.1016/j.jstrokecerebrovasdis.2010.05.002

27. Finlayson O, Kapral M, Hall R, Asllani E, Saposnik G, et al. Risk factors, inpatient care, and outcomes of pneumonia after ischemic stroke. Neurology. 2011; 77: 1338–1345. doi: 10.1212/WNL.0b013e31823152b1 PMID: 21940613

28. Walter U, Knoblich R, Steinhagen V, Donat M, Benecke R, Kloth A. Predictors of pneumonia in acute stroke patients admitted to a neurological intensive care unit. J Neurol. 2007; 254: 1323–1329. doi: 10.1007/s00415-007-0520-0 PMID: 17361338

29. Upadya A, Thorevska N, Sena KN, Manthous C, Arnouten-Adejegun Y. Predictors and consequences of pneumonia in critically ill patients with stroke. J Crit Care. 2004; 19: 16–22. PMID: 15101001

30. Yeh S-J, Huang K-Y, Wang T-G, Chen Y-C, Chen C-H, Tang S-C, et al. Dysphagia screening decreases pneumonia in acute stroke patients admitted to the stroke intensive care unit. J Neurol Sci. 2011; 306: 38–41. doi: 10.1016/j.jns.2011.04.001 PMID: 21549394

31. Doggett DL, Tappe KA, Mitchell MD, Chapell R, Coates V, Turkelson CM. Prevention of pneumonia in elderly stroke patients by systematic diagnosis and treatment of dysphagia: an evidence-based comprehensive analysis of the literature. Dysphagia. 2001; 16: 279–295. doi: 10.1007/s00455-001-0087-3 PMID: 11720404

32. Momosaki R, Abo M, Watanabe S, Kakuda W, Yamada N, Mochio K. Functional magnetic stimulation using a parabolic coil for dysphagia after stroke. Neuromodulation J Int Neuromodulation Soc. 2014; 17: 637–641; discussion 64. doi: 10.1111/ner.12137

33. Li L, Li Y, Huang R, Yin J, Chen Y, Shi J. The value of adding transcutaneous neuromuscular electrical stimulation (VitalStim) to traditional therapy for post-stroke dysphagia: a randomized controlled trial. Eur J Phys Rehabil Med. 2015; 51: 71–78. PMID: 25052012

34. Mann G, Hankey GJ, Cameron D. Swallowing function after stroke: prognosis and prognostic factors at 6 months. Stroke J Cereb Circ. 1999; 30: 744–748.
35. Heckert KD, Komaroff E, Adler U, Barrett AM. Postacute reevaluation may prevent Dysphagia-associated morbidity. Stroke J Cereb Circ. 2009; 40: 1381–1385. doi: 10.1161/STROKEAHA.108.533489

36. Langdon PC, Lee AH, Binns CW. Dysphagia in acute ischaemic stroke: severity, recovery and relationship to stroke subtype. J Clin Neurosci Off J Neurosurg Soc Australas. 2007; 14: 630–634. doi: 10.1016/j.jocn.2006.04.009

37. Lawrence ES, Coshall C, Dundas R, Stewart J, Rudd AG, Howard R, et al. Estimates of the prevalence of acute stroke impairments and disability in a multiethnic population. Stroke J Cereb Circ. 2001; 32: 1279–1284.

38. Veis SL, Logemann JA. Swallowing disorders in persons with cerebrovascular accident. Arch Phys Med Rehabil. 1985; 66: 372–375. PMID: 4004534

39. Hamdy S, Aziz Q, Rothwell JC, Crane R, Hughes D, Tallis RC, et al. Explaining oropharyngeal dysphagia after unilateral hemispheric stroke. Lancet Lond Engl. 1997; 350: 686–692. doi: 10.1016/S0140-6736(97)02068-0

40. De Marchis GM, Kohler A, Renz N, Arnold M, Mono M-L, Jung S, et al. Posterior versus anterior circulation strokes: comparison of clinical, radiological and outcome characteristics. J Neurol Neurosurg Psychiatry. 2011; 82: 33–37. doi: 10.1136/jnnp.2010.211151 PMID: 20802030

41. Sato S, Toyoda K, Uehara T, Toratani N, Yokota C, Moriwaki H, et al. Baseline NIH Stroke Scale Score predicting outcome in anterior and posterior circulation strokes. Neurology. 2008; 70: 2371–2377. doi: 10.1212/01.wnl.0000304346.14354.0b PMID: 18434640

42. Sarikaya H, Arnold M, Engelter ST, Lyrer PA, Mattle HP, Georgiadis D, et al. Outcomes of intravenous thrombolysis in posterior versus anterior circulation stroke. Stroke J Cereb Circ. 2011; 42: 2498–2502. doi: 10.1161/STROKEAHA.110.607614

43. Langdon PC, Lee AH, Binns CW. High incidence of respiratory infections in “nil by mouth” tube-fed acute ischemic stroke patients. Neuroepidemiology. 2009; 32: 107–113. doi: 10.1159/000177036 PMID: 19039243

44. Langmore SE, Terpenning MS, Schork A, Chen Y, Murray JT, Lopatin D, et al. Predictors of aspiration pneumonia: how important is dysphagia? Dysphagia. 1998; 13: 69–81. doi: 10.1007/PL00009559 PMID: 9513300

45. Langmore SE, Skarupski KA, Park PS, Fries BE. Predictors of aspiration pneumonia in nursing home residents. Dysphagia. 2002; 17: 298–307. doi: 10.1007/s00455-002-0072-5 PMID: 12355145

46. Mills B, Gosney M, Jack CIA, Martin MV, Wright AE. Acute stroke predisposes to oral gram-negative bacilli—a cause of aspiration pneumonia? Gerontology. 2003; 49: 173–176. doi: 69171 PMID: 12679608

47. Metheny NA, Chang Y-H, Ye JS, Edwards SJ, Defer J, Dahms TE, et al. Pepsin as a marker for pulmonary aspiration. Am J Crit Care Off Publ Am Assoc Crit-Care Nurses. 2002; 11: 150–154.

48. Brogan E, Langdon C, Brookes K, Budgen C, Blacker D. Respiratory infections in acute stroke: nasogastric tubes and immobility are stronger predictors than dysphagia. Dysphagia. 2014; 29: 340–345. doi: 10.1007/s00455-013-9514-5 PMID: 24445382

49. Fournier F, Cau-Pottier E, Boutigny H, Roussel-Delvallez M, Jourdain M, Chopin C. Effects of dental plaque antiseptic decontamination on bacterial colonization and nosocomial infections in critically ill patients. Intensive Care Med. 2000; 26: 1239–1247. PMID: 11089748

50. Drakulovic MB, Torres A, Bauer TT, Nicolas JM, Nogué S, Ferrer M. Supine body position as a risk factor for nosocomial pneumonia in mechanically ventilated patients: a randomised trial. Lancet Lond Engl. 1999; 354: 1851–1858. doi: 10.1016/S0140-6736(98)12251-1

51. Goldhill DR, Imhoff M, McLean B, Waldmann C. Rotational bed therapy to prevent and treat respiratory complications: a review and meta-analysis. Am J Crit Care Off Publ Am Assoc Crit-Care Nurses. 2007; 16: 50–61; quiz 62.

52. Daniels SK, Anderson JA, Willson PC. Valid items for screening dysphagia risk in patients with stroke: a systematic review. Stroke J Cereb Circ. 2012; 43: 892–897. doi: 10.1161/STROKEAHA.111.640946