Cerebral blood flow velocity changes during upright positioning in bed after acute stroke: an observational study

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ABSTRACT

Objectives: National guidelines recommend mobilisation in bed as early as possible after acute stroke. Little is known about the influence of upright positioning on real-time cerebral flow variables in patients with stroke. We aimed to assess whether cerebral blood flow velocity (CBFV) changes significantly after upright positioning in bed in the acute stroke phase.

Design: Observational study.

Participants: 47 patients with acute ischaemic stroke measured in the subacute phase after symptom onset and 20 healthy controls.

Primary and secondary outcome measures: We recorded postural changes in bilateral transcranial Doppler (primary outcome) and simultaneously recorded near-infrared spectroscopy, end-tidal CO2, non-invasive blood pressure data and changes in neurological status (secondary outcomes).

Methods: Postures included the supine, half sitting (45°), sitting (70°) and Trendelenburg (−15°) positions. Using multilevel analyses, we compared postural changes between hemispheres, outcome groups (using modified Rankin Scale) as well as between patients and healthy controls.

Results: The mean patient age was 62±15 years and median National Institute of Health Stroke Scale score on admission was 7 (IQR 5–14). Mean proportional CBFV changes on sitting were not significantly different between healthy controls and affected hemispheres in patients with stroke. No significant differences were found between affected and unaffected stroke hemispheres and between patients with unfavourable and favourable outcomes. During upright positioning, no neurological worsening or improvement was observed in any of the patients.

Conclusions: No indications were found that upright positioning in bed in mild to moderately affected patients with stroke compromises flow and (frontal)oxygenation significantly during the subacute phase of stroke. Supine or Trendelenburg positioning does not seem to augment real-time flow variables.

INTRODUCTION

Brain ischaemia is a potentially reversible process that is dependent on restoration of cerebral blood flow (CBF) within a time window of cellular viability that varies according to the severity and duration of flow...
cessation. Any activity that alters CBF during the acute phase of ischaemic stroke may, at least theoretically, directly contribute to extension of the penumbra to the ischaemic core and compromise clinical outcome. National guidelines recommend mobilisation in and out of bed as early as possible after stroke.2–4 Such early upright positioning prevents choking, cardiopulmonary deconditioning and may improve early plasticity of the brain.5 6 However, there is insufficient evidence in support of early upright positioning after acute stroke.7 Also, there is fear that mobilisation in the setting of a (potentially) disturbed cerebral autoregulation leads to gravity induced diminished CBF to the penumbra and, potentially, extension of the infarct core.8 As a result, early upright positioning is not universally accepted by stroke physicians.

Little is known in patients with acute stroke about the influence of different body positions on blood pressure (BP) or on real-time cerebral flow variables. Transcranial Doppler (TCD) may be used to assess CBF velocity (CBFV) at the stroke unit bedside. Nursing patients ‘heads down’ has been advocated, based on a 20% augmentation of CBFV when comparing supine with 30° head elevation in 20 patients severely affected with stroke.9 If such relatively minor changes in head position have such an impact, what would the consequences of more pronounced changes in position be in a representative sample of stroke unit patients?

In this study, we sought to assess the impact of the various head positions on CBFV comparing (1) hemispheres affected by stroke and unaffected hemispheres and (2) affected hemispheres and healthy controls. Any effect of upright positioning on neurological status and outcome was explored.

METHODS

Study design and patients

This observational study was conducted in the stroke unit of a teaching hospital between August 2008 and May 2011. Patients aged over 18 years with a first ever ischaemic stroke in the middle cerebral artery (MCA) territory were considered eligible. The measurements were taken on the stroke unit and mainly because of practical reasons (installation on stroke unit, informed consent approval and no interference with urgent clinical studies or consultations) performed in the subacute phase. Our protocol dedicated that the measurements had to be completed between 6 and 16 h after stroke symptom onset, mimicking the period of early in bed mobilisation initiation during daytime. Patients had to be conscious, cooperative, haemodynamically stable without fever and display flow in both M1 segments of the MCA. Moreover, patients deemed to be too severely affected by the stroke or too severely ill to be positioned upright (clinical/nursing judgement) were not included in this study. Brain CT, ECG, bilateral duplex carotid ultrasound and laboratory investigations were performed. Thrombolytic therapy was administered according to (inter)national guidelines. Use of any medication, especially antiarrhythmics and antihypertensive medication, was registered. The National Institutes of Health Stroke Scale score (NIHSS) was used to determine neurological deficit. The stroke subtype was classified using the Oxfordshire Community Stroke Project classification,10 and stroke aetiology according to Trial of Org 10172 in Acute Stroke Treatment (TOAST) definitions.11 The modified Rankin Scale (mRS) score at 3 months was determined by telephone interview by a certified assessor blinded to study data. Twenty healthy controls were recruited at random from (retired) work associates and their medical histories were reviewed. All experiments complied with the tenets of the Declaration of Helsinki and the study protocol was reviewed and approved by our institutional review board. Study participants provided written informed consent. Next of kin gave consent when participants were unable to provide it.

Sample size calculation

In the literature, SD of the intradividual supine to sitting MCA CBFV difference is estimated to be around ±5% for elderly hospitalised patients12 and ±20% for patients with severe stroke going from the 30° to 0° position.13 To detect a difference in mean CBFV of more than 6 cm/s (10–20%) between remaining supine and sitting upright in bed with a probability (power) of 90% when testing at a two-sided significance level of 5%, a sample size of 45 participants was required.

Procedures

BP was measured with a finger plethysmograph (Finometer-Pro, Finapress Medical Systems, Amsterdam, the Netherlands). The cuff was applied to the middle finger of the non-dominant or paretic hand placed at the heart level with the arm across the chest and supported by a sling. TCD was performed by the same investigator (MA) over the temporal bone window on both sides. MCA wave forms were identified at a depth range of 40–60 mm and a stable forward waveform with good intensity was selected for monitoring (Nicolet Pioneer TC8080, Carefusion Corporation, San Diego, California, USA). Patients with permanent MCA occlusion on TCD were excluded. TCD probes (2 MHz) were mounted by a head frame to ensure a constant angle of insonation throughout the positioning procedure. Bifrontal near-infrared spectroscopy (NIRS) recordings were added from 2010 onwards for regional cerebral oxygenation (rSO2, %) calculation. The INVOS 5100C NIRS device (Somanetics Corp, Detroit, Michigan, USA) has a sampling frequency of 0.2 Hz. Adhesive optodes were placed on each side of the forehead. End-tidal CO2 (EtCO2, kPA) was measured by a mask using an infrared CO2 analyser (Capnomac Ultima, GE Healthcare, Chalfont St Giles, UK) as an approximation of partial arterial CO2 pressure.
Signals
The signals of the BP and EtCO₂ waveforms, spectral envelope of MCA CBFV, and rSO₂ as well as the position marker signals were sampled at 250 Hz and synchronously captured using an A/D converter. Signals were recorded with LabVIEW software (LabVIEW V 7.1, National Instruments, Austin, Texas, USA) for online inspection and were stored for postprocessing. The MCA CBFV \( \text{mean} \) (cm/s) was calculated by using beat-to-beat mean values that were obtained by triggering off the ascending slope of the BP curve. Mean arterial pressure (MAP) at the MCA level (MAP\(_{\text{mca}}\), mm Hg) was automatically calculated from MAP measured at the heart level and the vertical finger-to-TCD probe distance. In the upright and Trendelenburg positions, great care was taken to secure the hand with the finger plethysmograph cuff at the heart level (approaching the right atrium level). The beginning and ending time points of the postural transitions were marked, but these data periods (±30 s) were not used for analysis.

Predefined positioning protocol
After admission to the stroke unit, patients were instrumented. Participants were requested to abstain from moving and talking in order to minimise artefacts and activity-related CBFV changes. Great care was taken to keep the head in the neutral midline position (without shifting of the headband) and the BP cuff at the heart level during positioning. A test run was performed. After 5 min of supine rest (baseline (eg, first) supine), the participants were passively tilted by changing the head of the bed end angles to the 45° (half sitting), 70° (sitting) and −15° (Trendelenburg) positions for 3–5 min each. Patients were brought in the sitting position twice. Periods of tilts were interspaced with 3–5 min of supine rest to re-establish control values (supine position).
Before and after establishing each position, the motor score of NIHSS (affected arm and leg) was evaluated. Control participants were recorded in a quiet, temperature-controlled research room using the same predefined positioning protocol.

Statistical analysis
We calculated a mean value for every position period. To compare responses in each position, percentage changes in the variables (MCA CBFV\(_{\text{mean}}\), rSO₂, MAP\(_{\text{mca}}\), heart rate (HR) and EtCO₂) were calculated as the differences between the positions after normalisation to the mean baseline (eg, first) supine value. Normalisation was carried out to be able to compare physiological responses between patients, healthy controls and hemispheres, where absolute individual values might differ from baseline. We averaged these percentage changes to obtain a mean percentage change for each of the four different positions: supine, half sitting, sitting and Trendelenburg. For control participants, the mean of the MCA CBFV\(_{\text{mean}}\) and rSO₂ signals was taken. To account for and to take advantage of the correlation between repeated measures in the same individual, a multilevel model was used to estimate the effects of positioning and affected hemisphere on changes in the main variables (MCA CBFV\(_{\text{mean}}\) and rSO₂).\(^{15}\) The relation between MCA CBFV\(_{\text{mean}}\) and outcome (dichotomised as mRS 0–2 for favourable outcome) was analysed in a similar way. Statistical analysis was performed with SAS (V.9.2). Significant carotid stenosis, stroke subtype or aetiology (TOAST criteria), NIHSS, age and sex were used as potential confounding covariates in stroke group comparisons.\(^{14}\) Age, sex and history of hypertension, cardiac disease and diabetes were used as covariates for healthy control and stroke group comparisons. Bonferroni adjustments for multiple comparisons were used for all results presented.

RESULTS
In eight patients with subacute stroke, TCD or BP investigation was unsuccessful. Eventually, we enrolled 47 patients (table 1). The mean patient age was 62 ±15 years and median baseline NIHSS was 7 points (IQR 5–14 points). At 3 months, 29 patients (62%) had a favourable outcome. NIRS recordings were performed in 32 patients (68%) and 16 controls (80%).
We compared the cerebral haemodynamic postural (MCA CBFV\(_{\text{mean}}\) and rSO₂) changes between positions, patients with stroke and healthy controls, and between affected and unaffected hemispheres. A complete overview of the comparisons is presented in online supplementary tables 1–3 (http://bmjopen.bmj.com).

Figure 1 shows that cerebral haemodynamic changes were most pronounced when moving from the supine to the sitting position (70°) for patients (affected hemisphere) and healthy controls (figure 1). The results presented below therefore focus on changes from the supine to the sitting position. The accompanying systemic haemodynamic (MAP\(_{\text{mca}}\) and HR) and respiratory responses (EtCO₂) are shown in figure 2.
In the sitting position for the affected and unaffected hemispheres, MCA CBFV\(_{\text{mean}}\) decreased in 28 patients (60%) and rSO₂ decreased in 18 patients (55%). None of the univariate results presented below were different after adjustment for predefined covariates, except for the outcome analysis.

Positional CBFV changes on sitting in patients with stroke
The mean MCA CBFV\(_{\text{mean}}\) decreased significantly with movement to the sitting position, (mean difference −4%, 95% CI −6% to −3%; \(p=0.005\)) for the affected hemisphere. For the unaffected hemisphere, this effect was not significant (mean difference −2%, 95% CI −5% to 0%; \(p=1.0\)).

Positional CBFV changes on sitting comparing patients with stroke and healthy controls
The mean MCA CBFV\(_{\text{mean}}\) changes on sitting were comparable between healthy controls and affected hemispheres of
patients with stroke (mean difference −4% for the affected hemisphere (95% CI −11% to 3%; p=1.0) as well as unaffected hemispheres (mean difference −5%, 95% CI −10% to 1%; p=0.18)).

Positional CBFV changes on sitting comparing hemispheres in patients with stroke

The mean MCA CBFV mean was not different between the affected and the unaffected hemispheres in the
sitting position (mean difference $-2\%$, 95% CI $-5\%$ to $2\%; p=1.0$).

**Positional changes of $r$SO$_2$, MAP$_{mca}$, HR and EtCO$_2$ on sitting**

The mean $r$SO$_2$ values decreased significantly with movement to the sitting position for the affected hemisphere (mean difference $-2\%$, 95% CI $-3\%$ to $-1\%; p=0.006$ and unaffected hemisphere (mean difference $-2\%$, 95% CI $-3\%$ to $-1\%; p=0.009$)). In agreement with the MCA CBFV$_{mean}$ results, the mean $r$SO$_2$ postural responses were not significantly different between both hemispheres and between patients and healthy controls. The CBFV$_{mean}$ and $r$SO$_2$ changes were positively correlated for the affected hemisphere ($r=0.33$, $p=0.01$).

In patients with stroke, MAP$_{mca}$ decreased on average by 15% in the sitting position (95% CI $-19\%$ to $-10\%; p<0.001$). This was similar to controls (17%, 95% CI $-24\%$ to $-11\%; p<0.001$; figure 2). MAP$_{mca}$ increased slightly during the course of the experiment for patients (mean 5%, 95% CI 2% to 7%) and healthy controls (mean 2%, 95% CI $-1\%$ to 6%). The postural responses were accompanied by only minor HR changes. EtCO$_2$ decreased on average by 3% (95% CI $-5\%$ to $-1\%; p<0.001$) for patients in the sitting position.

**Positional changes, neurological status and relation with outcome in patients with stroke**

We observed no neurological worsening or improvement during the postural changes in any of the patients. There were no significant differences between the mean MCA CBFV$_{mean}$ changes on upright and Trendelenburg positioning in patients with unfavourable compared to favourable outcome after correction for confounding covariates (mean difference on sitting, for the affected hemisphere: $-5\%$, 95% CI $-14\%$ to $5\%; p=1.0$).

**DISCUSSION**

Our study showed that following an ischaemic stroke in the MCA territory, early upright positioning in bed resulted in minor average changes in CBFV only in the subacute phase. The postural responses were not significantly different between the affected and unaffected hemispheres, and between patients and healthy controls. Passive upright positioning was not accompanied by worsening of the (motor) stroke severity score. No improvement was noticed in the supine and Trendelenburg positions. The relative postural haemodynamic and EtCO$_2$ changes were largely comparable between patients and controls. This study found no indications that upright positioning compromises flow or (frontal) oxygenation significantly in the affected MCA territory in mild to moderately affected patients with acute stroke. No association was found between flow changes and functional outcome. Our results suggest effective cerebral and/or systemic haemodynamic adjustments to overcome gravitational force during postural changes in the subacute phase of mild to moderate stroke. The potential of the Trendelenburg position to augment CBF seems to be limited and may not outweigh concurrent cardiopulmonary complications.

Although TCD measurements cannot be used to calculate CBF, the observed relative CBFV changes can be considered proportionate to CBF changes as long as the angle of insonation and the MCA diameter remain constant during the brief test interval. The MCA diameter is thought to remain constant over a ±30 mm Hg range of BP and ±2 kPa of CO$_2$.

This requirement was fulfilled in our study. TCD is prone to several errors because the measurements are operator dependent and patient dependent. Even small probe position changes may alter the CBFV readings. However, frontal cerebral oxygenation ($r$SO$_2$) followed CBFV changes in size and direction in the affected hemisphere, which further strengthens our findings. Assuming a constant arterial O$_2$ content and cerebral metabolic rate during the measurement, $r$SO$_2$ is predominantly thought to remain constant over a ±30 mm Hg range of BP and ±2 kPa of CO$_2$. Furthermore, TCD monitoring is the only non-invasive method to monitor CBF changes continuously at the bedside. It is generally accepted that CBF changes are closely reflected by TCD readings.

Patients in our study predominantly had mild to moderate strokes. Patients with MCA occlusion were excluded. It was left to clinical and nursing judgement whether patients were deemed too severely affected by the stroke or too severely ill to be positioned upright for this study. Although NIHSS was not found to be a confounder for postural CBFV changes, too few patients with severe stroke (NIHSS>16) were included to be able to generalise our findings to patients with severe strokes. Furthermore, fully ambulatory patients may have been less likely to be included. The practical implication of the latter drawback in generalisation is probably limited. When mild to moderate strokes are unlikely to be affected by upright positioning, very mild strokes are even less likely to be affected.
Three studies have investigated the postural effects on CBFV in acute stroke.9 10 20 Wojner et al found on average a 20% CBFV increase in the supine position in 20 patients with an acute MCA occlusion. However, a wide range of individual flow improvements was present (5–96% from baseline), with no data from the unaffected hemisphere being available.9 A small study with eight participants found important postural flow changes in four severely affected patients with incomplete MCA recanalisation on 24 h follow-up imaging (affected hemisphere).19 A more than 15% higher MCA flow was found in the supine position in 18 patients with large stroke with intracranial pressure (ICP) monitoring in the intensive care unit. This effect was only present in the affected hemisphere.20 Different results were obtained in a single-photon emission CT (SPECT) stroke study. Seven of eight patients with subacute stroke had a clear visual improvement in SPECT perfusion to the affected frontal, temporal or parietal lobes in the semirecumbent (30–45°), compared to the supine position. No differences were found in BP in both positions. Unfortunately, no demographics, clinical effect or respiratory changes of the more upright position were reported.21

Our study did not show relevant postural changes despite the larger position changes (70° instead of 30°). This is probably largely due to our predominant focus on mild to moderately affected patients with stroke. Also, in our study, changes in all variables were related to the individual baseline supine level (by normalisation), reducing the effects of large variations of measured parameters and difficulties with standardisation of the used techniques (TCD, NIRS and BP). In none of the four studies were flow data recorded simultaneously with NIRS and EtCO2. In the study of Hargroves et al, cerebral oxygenation as measured by NIRS in the first seven days after stroke was lowest in the upright position and highest in the supine position (difference estimated around 2–6%). These changes were also present in the unaffected hemisphere, although on visual inspection they were less marked.22 Furthermore, there might be an effect of anaesthesia, vasopressors, hypothermia and decompressive cranietomy on cerebral vasoregulation used in the intensive care stroke positioning study.23 Our results are in line with patients with subarachnoid haemorrhage in whom CBFV remained constant in the supine and 45° patient body positions, regardless of the development of vasospasm.24

It is widely believed that a sudden BP reduction can affect CBF and render elderly participants or patients with cardiovascular disease vulnerable to orthostatic symptoms.25 The sitting position might be dangerous, especially with globally impaired cerebral autoregulation, because of the susceptibility of the penumbral regions to local hypoperfusion in the acute and subacute phases. However, our physiological results indicate that sitting is probably well tolerated in mild to moderately affected patients with stroke. In 2008, the ‘Very Early Rehabilitation trail’ showed that very early out of bed mobilisation was safe and feasible in patients with stroke.26 This may be due to the counterbalancing effects of BP elevation, ICP reduction and increased venous return to stabilise cerebral perfusion pressure in the subacute phase. There is evidence for increased sympathetic activation in patients with acute stroke.27–29 Replication of this study should be considered in studying the (hyper)acute phase, longer sitting periods and using more sophisticated and direct methods that measure CBF regionally and estimate cerebral oxygen metabolism.30 Inclusion of more severely disabled patients (NIHSS around 10) would be preferable to being able to compare upright cerebral haemodynamics with the results from the ongoing ‘early mobilisation’ trails.26 Further research should also include more detailed (follow-up) information from extracranial and intracranial vessels pathology (using validated radiological or ultrasound criteria), recanalisation (proximal and distal), penumbra configuration and/or the presence of collaters.31–34 Follow-up studies several days after stroke may be warranted because local and systemic compensatory mechanisms may become exhausted, but around that time influence on penumbral salvage may be limited.35 Our results do not hold for sitting to the standing mobilisation, which may provoke different (probably larger) haemodynamic responses.36

In the paucity of results for the early out of bed mobilisation stroke trails,37 we suggest continuation of early upright positioning in patients with stroke in the stroke unit in the subacute phase and propose that continuous CBF measurements during early activity out of bed (standing) could be helpful in assuring safe and early mobilisation in the stroke unit.36

CONCLUSION
In conclusion, upright positioning in the stroke unit of mildly to moderately affected patients with stroke does not give significant reduction of CBF and (frontal) cerebral oxygenation surrogates during the subacute phase on the stroke unit. Supine or Trendelenburg positioning seems not to augment MCA CBFVmean or improve neurological status.

Contributors MJA, JWE and PV contributed to design and conceptualisation of the study, interpretation of the data, and drafting and revision of the manuscript. RS and JDK contributed to the interpretation of the data and revision of the manuscript. BK was responsible for the revision of the manuscript.

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Competing interests None.

Patient consent Obtained.

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Data sharing statement The authors are willing to share the data, as well as the additional unpublished data of follow-up studies (day 3 after stroke) and dynamic autoregulation results.
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