Remote Ischemic Limb Preconditioning After Subarachnoid Hemorrhage
A Phase Ib Study of Safety and Feasibility

Sebastian Koch, MD; Michael Katsnelson, MD; Chuanhui Dong, PhD; Miguel Perez-Pinzon, PhD

Background and Purpose—Making a limb transiently ischemic has been shown to induce ischemic tolerance in a distant organ. This phenomenon is known as remote ischemic limb preconditioning. We conducted a Phase IB study of remote ischemic limb preconditioning to determine the safety and feasibility of increasing durations of limb ischemia in patients with subarachnoid hemorrhage.

Methods—Patients with aneurysmal subarachnoid hemorrhage underwent limb preconditioning every 24 to 48 hours for 14 days. Limb preconditioning consisted of 3 5-minute inflations of a blood pressure cuff to 200 mm Hg around a limb followed by 5 minutes of reperfusion. In the lead-in phase, we preconditioned the upper extremities, but this proved impractical and we began preconditioning the leg in a similar manner. Ischemia times were then escalated to 7.5 and 10 minutes. After each session, a visual analog scale was obtained and the extremity examined for neurovascular complications.

Results—A total of 33 patients completed the study. Mean age was 53±12 years and mean Hunt Hess score was 2.4±0.9. In the lead-in phase, an average of 7.7±2.4 preconditioning sessions was completed with mean visual analog scale 3.6±3.4. In the dose escalation phase, an average of 8.6±2.1 preconditioning sessions was done with mean visual analog scale 1.8±2.2 and 2.5±2.9 for the 7.5- and 10-minute cohorts, respectively. No session was prematurely terminated due to subject discomfort. No objective signs of neurovascular injury were observed.

Conclusions—We found limb preconditioning to be safe and well tolerated, even at ischemia times of 10 minutes, in critically ill patients with subarachnoid hemorrhage. (Stroke. 2011;42:1387-1391.)

Key Words: delayed cerebral ischemia ■ limb preconditioning ■ remote ischemic preconditioning

Transient exposures to mild ischemia render the brain resistant to subsequently more severe ischemic injury. This is known as ischemic preconditioning and is one of the most powerful anti-ischemic strategies in laboratory models. Similarly, inducing ischemia in one organ also leads to ischemic tolerance in a different, distant organ. A simple example of this phenomenon, referred to as remote ischemic limb preconditioning, is that animals undergoing brief limb ischemia have smaller infarcts than animals not exposed to this intervention before stroke.1,2 Preconditioning induces multiple endogenous anti-ischemia defense mechanisms. These include an upregulation of nitric oxide, improved cerebral blood flow in the ischemic penumbrae, and down-regulation of inflammation and glutaminergic excitotoxicity.3,4 With ischemic preconditioning, cellular energy consumption is optimized and becomes more efficient, rendering the cell more resistant to ischemia.5–8

Only a few ischemic preconditioning trials have been completed. Brief cycles of arm ischemia before cardiovascular surgery or percutaneous cardiac intervention resulted in a reduction of periprocedural myocardial infarction, and fewer strokes.9–12 However, the therapeutic potential of preconditioning has remained largely unexplored in cerebrovascular disease. Only one randomized trial assessed the effect of two 10-minute cycles of leg ischemia before carotid endarterectomy. The intervention resulted in a statistically nonsignificant reduction of saccadic latency, a surrogate marker for hemispheric ischemia.13

Clinical settings such as carotid revascularization may be very suitable for a study of preconditioning. The risk of cerebral ischemia occurs over a short, well-defined period of time, allowing a preconditioning response to be elicited. Other settings frequently proposed for preconditioning include cardiac bypass surgery or subarachnoid hemorrhage with the ensuing risk of delayed cerebral ischemia (DCI).5,14,15 The high morbidity and mortality associated with DCI has made the investigation of innovative treatment approaches a priority in subarachnoid hemorrhage research.

We report on our preliminary experience with ischemic limb preconditioning after subarachnoid hemorrhage. The scientific rational for such a study is to determine if daily limb preconditioning after subarachnoid hemorrhage induces is-
chemic tolerance and protects from symptomatic vasospasm and DCI. Because the optimal duration of limb ischemia for preconditioning and the safety and tolerability of daily repetitive limb preconditioning in predominantly conscious but critically ill patients is uncertain, we aimed to establish both of these parameters in a preliminary trial of safety and feasibility. The purpose of this particular study was to assess the safety and feasibility of repetitive limb preconditioning and increasing durations of limb ischemia in patients with recent subarachnoid hemorrhage.

**Methods**

The study was conducted at a large community-based teaching hospital, which serves as a referral center for patients with subarachnoid hemorrhage. We enrolled subjects with acute subarachnoid hemorrhage due to a ruptured cerebral aneurysm, secured by either coiling or surgical clipping, within 96 hours from symptom onset.

Subjects with the following were excluded: (1) impaired prehospitalization functional status defined as modified Rankin Scale score >2; (2) subarachnoid hemorrhage from a lesion other than cerebral aneurysm; (3) Hunt Hess scale ≥4 at presentation; (4) history of peripheral vascular disease and ankle brachial index <0.7; (5) any extremity soft tissue, orthopedic, or vascular injury which in the clinical judgment of the investigator may pose a contraindication to preconditioning, for example, superficial wounds, cellulitis, fracture, deep vein thrombosis; (6) patients with symptomatic vasospasm at the time of enrollment; (7) neurogenic pulmonary edema or cardiac failure requiring inotropic support; or (8) severe or unstable concomitant chronic condition or disease.

All patients received standard treatment for subarachnoid hemorrhage, cerebral vasospasm, and DCI. In addition, all subjects underwent limb preconditioning throughout the risk period for cerebral vasospasm.

**Limb Preconditioning Protocol**

Subjects underwent a limb preconditioning session every 24 to 48 hours from the time of enrollment to Day 14 (or discharge if before Day 14). We initially induced ischemia in the upper extremity according to a method shown to protect the myocardium during cardiac surgery. A blood pressure cuff was placed over the arm and inflated to 200 mm Hg (or 20 mm Hg above systolic pressure if systolic pressure >200 mm Hg). The cuff remained inflated for 5 minutes followed by 5 minutes of reperfusion. Each preconditioning session consisted of 3 5-minute cycles of ischemia/reperfusion. We also included a control group who underwent a sham preconditioning procedure. Sham preconditioning consisted of minimal inflation to systolic pressure 30 mm Hg, just enough to cause some pressure on the extremity. Subjects were initially randomly assigned to true or sham preconditioning. After completion of each preconditioning session, subjects were asked (if possible) to rate pain on a visual analog pain scale (VAS), which ranges from 0 (no pain) to 10 (worst imaginable pain). After each session, the extremity was examined for tissue breakdown or signs of neurovascular injury.

**Statistical Analysis**

Statistical analyses include summary statistics presented as mean and SD for continuous variables and frequency and percentages for categorical variables. Between-group comparisons were made with analysis of repeated measures with significance set as P<0.05. Postanalysis of variance was done with Bonferroni multiple comparisons tests. Categorical variables were compared by means of a χ² test or Fisher exact test as appropriate. Analysis was performed with SPSS Version 17 (Chicago, IL).

**Results**

A total 34 patients were enrolled. One subject refused continued study participation due to headaches. The baseline characteristics of the 34 subjects are outlined in Table 1. The most frequent site of aneurysm was the anterior communicating artery (46%) followed by the posterior communicating artery (21%) and middle cerebral artery (12%). During the course of the study, mild transcranial Doppler vasospasm was
found in 22 of 33 (67%). Severe transcranial Doppler vaso-
spasm was diagnosed in 8 of 33 (24%) and DCI in 6 of 33
(18%). Two subjects (6%) died during the course of the study.
The mean National Institutes of Health Stroke Scale score at
14 days was 1.7±2.6. The mean 3-month modified Rankin
Scale score was 1.3±1.4.

**Lead-In Phase**

In our initial experience, we enrolled 21 subjects. The first 6
subjects (including 3 with sham preconditioning) underwent
arm preconditioning. Arm preconditioning proved impracti-
cal because monitoring devices and intravenous lines inter-
fereed with our ability to induce limb ischemia. The next 15
subjects (including 4 with sham procedure) underwent leg
preconditioning, which proceeded without difficulties. Those
subjects had on average 7.7±2.4 preconditioning sessions. In
full treatment patients who could respond, the mean VAS per
patient session was 3.6±3.4. For sham-conditioned patients,
the VAS was 0. No session was prematurely terminated due
to subject discomfort. No objective signs of neurovascular
injury were observed with no subjects experiencing skin
breakdown, prolonged discoloration, or temperature or pulse
disparities in the treated limb. Among the 21 patients, 9 lower extremity Duplex studies were obtained.

Deep vein thrombosis was found in 3; 2 were symptomatic,
in a patient undergoing arm preconditioning and 1 in the
leg of a patient who had the contralateral leg preconditioned.
These were felt to be unrelated to the intervention. An
asymptomatic partial occlusion occurred in 1 patient in the
preconditioned leg.

**Dose Escalation Phase**

We then escalated limb ischemia times to 7.5 and 10 minutes,
respectively, in cohorts of 6 subjects in each tier. Those
subjects underwent an average of 8.6±2.1 preconditioning
sessions. In those able to respond, the VAS was 1.8±2.2 and
2.5±2.9 for the 7.5- and 10-minute cohorts, respectively. The
VAS was higher in the 5-minute ischemia group, but this did
not differ significantly from the other 2 groups. No session
was prematurely terminated due to subject discomfort. No
objective signs of neurovascular injury were observed. There
were no cases of deep vein thrombosis.

**Outcome Data**

The main focus of our study was safety and subject tolera-
bility of the procedure, but we explored a potential efficacy
signal. Table 2 shows stroke severity and transcranial Doppler
data among the 4 groups. Given the small sample size,
imbalances between the groups due to chance are expected
and were found. Although none of the group differences were
statistically significant, the 10-minute limb ischemia group
was older, more likely to have hypertension, and had a higher
modified Rankin Scale score at 3 months. In that group, the
3-month outcome data were influenced by a smaller sample
size because 1 patient could not be contacted (the only loss of
follow-up in the study) and by the death of 1 subject (81 years
old). Excluding that outlier, the 3-month modified Rankin
Scale score (n=4) was 1.5±1.

**Discussion**

Ischemic preconditioning was first demonstrated in the myo-
cardium where transient occlusions of a canine coronary
artery resulted in a decrease in eventual myocardial infarct
size after permanent coronary occlusion.19 It is now well
recognized to confer ischemic protection across many differ-
tent organs and species.20,21 In the myocardium, ischemic
preconditioning remains one of the most powerful protective
strategies with reductions in infarct size between 50% and
70%.22

Similar tolerance to ischemia is also achieved in the brain.

Animals pretreated with 2 minutes of global ischemia showed
marked protection against CA1 neuronal loss after 5 minutes
of global ischemia.23 Simulating human stroke more closely
are mouse models that show brief occlusions of the middle
cerebral artery decrease final infarct size after temporary or
permanent middle cerebral artery occlusion.24 Cross-
tolerance is also observed between different organs. Pretreat-
ment with 15-minute bilateral forelimb ischemia resulted in a
54% increase in normal-appearing CA1 neurons in compari-
son to an untreated control group in an asphyxial cardiac
arrest model.25

Rendering a limb transiently ischemic to protect a distant
organ such as the brain is of great clinical importance because
this can be readily and noninvasively achieved in humans. A
study of 82 patients undergoing abdominal aneurysm repair
showed that 2 10-minute cycles of intraoperative crossclamp-
ing of the iliac artery led to a 27% reduction in troponin, 22%
a decrease in myocardial infarction, and 23% reduction in
renal impairment.12 In 2 studies of patients undergoing

coronary artery bypass surgery, myocardial injury was re-
duced in patients who underwent 3 cycles of arm ischemia,
induced with a blood pressure cuff, just before surgery.10,26

Perioperative troponin levels were decreased by 43% in
patients preconditioned in this manner.10 In 242 subjects, 3
5-minute cycles of arm ischemia before percutaneous cardiac
intervention has been associated with reduced troponin re-

<table>
<thead>
<tr>
<th>Limb Ischemia Duration</th>
<th>Age, Years</th>
<th>Hypertension, No. (%)</th>
<th>Hunt Hess</th>
<th>TCD Spasm Mild, No. (%)</th>
<th>TCD Spasm Severe, No. (%)</th>
<th>DCI, No. (%)</th>
<th>3-Month mRS Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 minute, n=7</td>
<td>49±9</td>
<td>1 (14)</td>
<td>2.7±0.5</td>
<td>4 (57)</td>
<td>0 (0%)</td>
<td>1 (14)</td>
<td>0.7±1.0</td>
</tr>
<tr>
<td>5 minutes, n=14</td>
<td>53±14</td>
<td>6 (43)</td>
<td>2.4±1.0</td>
<td>9 (64)</td>
<td>6 (43)</td>
<td>4 (29)</td>
<td>1.6±1.8</td>
</tr>
<tr>
<td>7.5 minutes, n=6</td>
<td>47±12</td>
<td>2 (33)</td>
<td>2.5±1.0</td>
<td>5 (83)</td>
<td>2 (33)</td>
<td>1 (17)</td>
<td>1.5±1.0</td>
</tr>
<tr>
<td>10 minutes, n=6</td>
<td>62±10</td>
<td>5 (83)</td>
<td>2.2±1.2</td>
<td>4 (67)</td>
<td>0 (0%)</td>
<td>0 (0)</td>
<td>2.5±1.7</td>
</tr>
</tbody>
</table>

TCD indicates transcranial Doppler; DCI, delayed cerebral ischemia; mRS, modified Rankin Scale.
lease, less electrocardiographic ST elevation, and fewer major adverse cardiovascular events. However, in a recent study with a larger population, a protective effect of remote ischemic preconditioning before coronary surgery could not be confirmed.27

Only a few clinical reports have assessed the preconditioning potential of the brain. In a small study of 12 patients undergoing cerebral aneurysm clipping for subarachnoid hemorrhage, direct preconditioning by temporary occlusion of a cerebral artery attenuated tissue hypoxia during subsequent prolonged arterial occlusion.28 A small case series reported that during carotid stenting, patients who became symptomatic at the time of carotid occlusion were able to tolerate subsequent occlusions during the procedure without any recurrent symptoms.29 In 70 patients randomized to 2 10-minute cycles of leg ischemia or sham preconditioning before endarterectomy, there was a nonsignificant reduction of saccadic latency (a surrogate for hemispheric injury) in patients who were preconditioned (32% versus 53%).30

Several questions remain regarding the best method to induce a preconditioning response. The optimal duration of limb ischemia remains uncertain. Is preconditioning a leg better than preconditioning an arm? The potentially larger volume of tissue affected by preconditioning a leg may translate into a greater biological effect. Clinical studies in cardiac medicine have examined 5-minute cycles of arm ischemia.10,11,26 However, it is not clear if this is the optimal method to induce a preconditioning response. In laboratory studies, limb ischemia times have typically exceeded 5 minutes (in the range of 10 to 30 minutes) and the hind leg of the animal was preconditioned.1,2,5,30–32

These questions led us to report our preliminary experience with limb preconditioning for neuroprotection. We were particularly concerned if repetitive and longer limb ischemia times, approaching durations achieved in laboratory animals, would be tolerated safely by patients with subarachnoid hemorrhage. Previous studies of limb preconditioning induced limb ischemia immediately before surgery with subjects already under general anesthesia. Our patient population was predominantly conscious, which raised additional concerns about potential tolerability. Our experience strongly suggests that this is the case. We did not find a single case of objective neurovascular injury and at no time did a subject refuse continued participation due to pain or discomfort.

In conclusion, we report the first clinical experience with remote ischemic preconditioning after subarachnoid hemorrhage. We found that a trial of repetitive ischemic limb preconditioning is feasible, safe, and well tolerated in awake patients with subarachnoid hemorrhage. The safety of limb preconditioning at ischemia times of at least 10 minutes needs further assessment in larger clinical trials.

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Disclosures
None.

References


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