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Is There Benefit from Stenting on Cognitive Function in Intracranial Atherosclerosis?

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Abstract

Background—Revascularization of stenotic cerebral arteries is hypothesized to improve cognition by increasing cerebral perfusion.

Aims—We compared cognition impairment among patients treated with percutaneous angioplasty and stenting (PTAS) and aggressive medical management (AMM) vs. AMM alone in the SAMMPRIS Trial.

Methods—In SAMMPRIS, 451 patients with recent TIA or stroke attributed to 70%-99% intracranial stenosis were randomized to PTAS plus AMM or AMM alone. Patients with stroke as the qualifying event with an NIHSS indicating aphasia or neglect were excluded from these analyses. Patients with a cerebrovascular event (ischemic stroke, cerebral infarct with temporary signs (CITS), or ICH) during follow-up were excluded from follow-up visit analyses. The Montreal Cognitive Assessment (MoCA) score was used to assess cognition impairment at
baseline, 4 months, 12 months, and closeout. Cognitive impairment was defined as MoCA <26. Mean MoCA scores and the percentage of patients with cognitive impairment were compared between treatment groups at each time point using t-tests and Chi-square tests. Differences in MoCA means from baseline to follow-up time points were compared using mixed model repeated measures ANOVA and Tukey-Kramer tests.

**Results**—There were no significant differences between the treatment groups for mean MoCA at any time point. Mean MoCA scores improved in both groups. The percentage of patients with cognitive impairment in the AMM vs. PTAS groups was not significantly different at any time point.

**Conclusions**—Revascularization with PTAS showed no improvement in cognitive impairment over aggressive medical management alone among patients who did not have recurrent cerebrovascular events during follow-up.

**Keywords**
intracranial stenosis; stroke; cognition; revascularization; angioplasty and stenting

**Introduction**
Cognitive function is an important secondary outcome in stroke prevention studies, particularly in trials that study interventions affecting cerebral blood flow. Revascularization procedures of stenotic cerebral or pre-cerebral arteries are of particular interest due to a hypothesized improvement in brain perfusion resulting in preservation or improvement of cognition, independent of stroke outcomes. For this reason, cognitive impairment was a pre-specified secondary outcome in the Stenting versus Aggressive Medical Therapy for Intracranial Arterial Stenosis (SAMMPRIS) trial that studied revascularization of stenotic cerebral arteries. While SAMMPRIS showed that aggressive medical management (AMM) alone was superior to percutaneous angioplasty and stenting (PTAS) plus AMM for secondary stroke prevention\(^1\), we sought to determine if patients treated with PTAS would have less cognitive impairment, independent of stroke recurrence, compared to AMM alone.

**Methods**
The design, participant characteristics and primary outcome results of SAMMPRIS have been previously reported.\(^{1-3}\) In brief, patients with recent transient ischemic attack (TIA) or stroke attributed to 70-99% intracranial stenosis were randomized to treatment with PTAS plus AMM or AMM alone. Patients randomized to PTAS were required to undergo PTAS within 3 business days of randomization. In order to minimize the effect of focal neurological deficits from stroke (e.g. aphasia or neglect) on cognitive function, patients presenting with stroke as the qualifying event who had a National Institutes of Health Stroke Scale (NIHSS) score that included points for aphasia or neglect were excluded from these analyses. Similarly, in order to assess the impact of treatment on cognitive function, independent of the treatment's stroke prevention impact, patients with symptomatic cerebrovascular events (ischemic stroke, cerebral infarct with temporary signs, or intracerebral hemorrhage) during follow-up were excluded from these analyses. Details
regarding the adjudication of events are described elsewhere\(^2\). In brief, study neurologists who were not masked to treatment assignment evaluated patients with a potential event. Patients with neurological events that were potentially difficult to classify (a TIA lasting > 1 hour or mild ischemic stroke [an increase in the patient's National Institute of Health Stroke Scale of < 4 from study entry]) were evaluated by a second neurologist who was masked to treatment. Both neurologists’ assessments were sent for central adjudication by neurology adjudicators who were masked to treatment assignment.

The Montreal Cognitive Assessment (MoCA) was used to assess cognitive impairment. MoCA is a screening tool that is sensitive to cognitive deficits arising from stroke and vascular cognitive impairment\(^4\). MoCA was performed at baseline, 4 months, 12 months, and closeout. Cognitive impairment was defined as a MoCA score < 26, the standard value that has 90% sensitivity for detecting mild cognitive impairment\(^5\). Additional analyses also examined cognitive impairment using a MoCA < 23, and mean MoCA scores. We also examined a MoCA sub-score of 11 points that focused on frontal and subcortical function (MoCA FS), which used the Trails B, clock drawing, attention (3 tasks), and fluency portions of the MoCA. In addition, the percentage of patients in each treatment group who had a change in MoCA of 1 or more points was also compared for the group overall and the subset of patients with more than 3 years of follow-up.

Mean MoCA scores and the percentage of patients with cognitive impairment using the MoCA thresholds and FS sub score described above were compared between treatment groups at each time point using t-tests and Chi-square tests. The percentage of patients with a change in MoCA +/- 1 point was compared between treatment groups using Chi-square tests. Differences in MoCA means from baseline to follow-up time points for each treatment group were compared using mixed model repeated measures, ANOVA, and Tukey-Kramer tests and adjusted for age, gender, hypertension, diabetes, lipid disorder, old infarcts, stroke as the qualifying event, and location of the symptomatic artery.

### Results

Of the 451 patients enrolled in SAMMPRIS, 371 had MoCA recorded at baseline and did not have a NIHSS that indicated aphasia or neglect (n=188 in AMM group, n=183 in PTAS group). At 12 months, 253 of these patients had MoCA data and did not have a cerebrovascular event in the preceding 12 months. At closeout, 199 patients had MoCA data and did not have a cerebrovascular event prior to closeout. Mean follow-up was 3.05 and 3.12 years in the AMM and PTAS groups, respectively. The number of subjects with at least 3 years of follow-up were similar between the AMM (n=59) and PTAS (n=54) groups.

The percentage of patients with cognitive impairment (MoCA < 26) in the AMM vs PTAS groups was not significantly different at baseline (53 vs 56%, \(p=0.55\)), 12 months (42 vs 40%, \(p=0.70\)) and closeout (43 vs 38%, \(p=0.48\)). Similarly, at both 12 months and closeout, cognitive function was not impacted by treatment when using a MoCA cutoff of < 23 (\(p=0.94\) and \(p=0.81\), respectively). Mean MoCA scores were not different between AMM vs. PTAS at 12 months (25.4 ± 3.73 vs. 25.65± 3.89, \(p=0.55\)) and closeout (25.58±3.61 vs. 25.66±3.76, \(p=0.90\)). There was also no difference in mean MoCA FS at 12 months.
(p=0.93) and closeout (p=0.69). The percentage of patients in each treatment group who had an increase or decrease of MoCA score from baseline to closeout was not different for the group overall (p=0.59) or for only subjects with more than 3 years of follow-up (p=0.65), as shown in Tables 1 and 2. Patients in both treatment groups had a statistically significant (p<0.05) approximately 1 point increase in MoCA score from baseline to the follow-up visits, as shown in Figure 1. Treatment assignment was not associated with a change in MoCA score from baseline to 1 year or closeout (p= 0.4747 and p=0.4706, respectively), even after adjusting for age, gender, hypertension, diabetes, lipid disorder, old infarcts, stroke as the qualifying event, and location of the symptomatic artery (p=0.4656 and p=0.5640).

**Discussion**

In SAMMPRIS, revascularization with PTAS did not provide improvement in cognitive assessment scores over aggressive medical management alone among patients who did not have recurrent cerebrovascular events during follow-up. While it has been hypothesized that reduced perfusion through a large cerebral artery may lead to cognitive decline and therefore reperfusion of stenotic cerebral arteries would improve cognition, there is a paucity of data that supports this hypothesis. Few randomized trials studying revascularization in patients with large artery cerebrovascular disease have assessed cognition and the available data are limited to only extracranial arteries\(^6\)-\(^8\), not intracranial arteries. Both the Asymptomatic Carotid Artery Study (ACAS), which studied carotid endarterectomy versus medical therapy, and the Randomized Evaluation of Carotid Occlusion and Neurocognition (RECON) trial (an ancillary study of the Carotid Occlusion Surgery Study (COSS) that studied extracranial–intracranial bypass for carotid occlusion), showed no benefit of revascularization on cognition over time\(^6\),\(^7\). ACAS reported a non-significant decline in the MMSE in both treatment groups during 5 years of follow-up\(^6\). RECON showed no difference in the change in cognitive function over time between medical or revascularization groups during 2 years of follow-up using a battery of 14 neurocognitive tests.\(^7\) Together, analyses of these 3 randomized clinical trials provide no support for the hypothesis that revascularization of large cerebral arteries provides any improvement in cognitive assessment scores.

Overall, SAMMPRIS patients in both treatment groups did have a small improvement in performance in MoCA testing during a mean follow-up of 3 years. However, both groups also had a large percentage of patients with at least minimal decline. Given that multimodal risk factor control, including healthy diet, exercise, and vascular risk factor medications, improves cognitive outcome in healthy older adults,\(^9\) it is possible that the intensive risk factor treatment provided in SAMMPRIS\(^3\) may have resulted in some cognitive improvement in this high-risk stroke population. Future analyses on the impact of risk factor control on MoCA scores in SAMMPRIS may address this issue. However, the increase in MoCA scores seen in both groups during follow-up may also be due to a possible learning effect of the MoCA, as increases of 0.9 points at 1 month have been reported\(^5\).

A limitation of our study is that we cannot account for the impact of silent cerebral infarcts, which may have been more prevalent in one treatment group, on the MoCA scores because brain imaging was not required in patients without neurological symptoms. Similarly,
angiography during follow-up was not permitted, so the impact of restenosis or occlusion on MoCA scores cannot be assessed. The use of MoCA to assess cognitive function may also be considered a limitation. While the MoCA can be sensitive for screening for cognitive impairment, it does not provide the detailed assessment of subtle changes in cognitive function that a more comprehensive neuropsychological evaluation would. Future studies of treatments for symptomatic intracranial stenosis could address this issue by performing more detailed neuropsychiatric testing. However, given that stenting of stenotic intracranial arteries is not recommended for stroke prevention\(^\text{10}\), future studies to assess the impact of revascularization on stroke prevention and/or cognition are likely to be limited to very select high-risk patients and not generalizable to all patients with intracranial stenosis.

**Summary**

In conclusion, among patients with symptomatic severe intracranial atherosclerotic stenosis, revascularization does not provide any independent benefit on cognitive assessment scores over medical therapy and therefore is unwarranted for that purpose.

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**REFERENCES**


Figure 1.
MoCA Score by Treatment Assignment During Follow-up
### Table 1
Change in MoCA score among SAMMPRIS patients by treatment assignment

<table>
<thead>
<tr>
<th>Change in MoCA points</th>
<th>AMM (Medical therapy only)</th>
<th>PTAS (stenting and medical therapy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increase &gt;= 1</td>
<td>48 (37.5%)</td>
<td>39 (32.0%)</td>
</tr>
<tr>
<td>No change</td>
<td>19 (14.8%)</td>
<td>20 (16.4%)</td>
</tr>
<tr>
<td>Decrease &gt;= 1</td>
<td>61 (47.7%)</td>
<td>63 (51.6%)</td>
</tr>
</tbody>
</table>

Chi-square p-value = 0.655
Table 2
Change in MoCA score among SAMMPRIS patients with > 3 years of follow-up by treatment assignment

<table>
<thead>
<tr>
<th>Change in MoCA points</th>
<th>AMM (Medical therapy only)</th>
<th>PTAS (stenting and medical therapy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increase &gt;= 1</td>
<td>20 (33.9%)</td>
<td>18 (33.3%)</td>
</tr>
<tr>
<td>No change</td>
<td>7 (11.8%)</td>
<td>10 (18.5%)</td>
</tr>
<tr>
<td>Decrease &gt;= 1</td>
<td>32 (54.2%)</td>
<td>26 (48.2%)</td>
</tr>
</tbody>
</table>

Chi-square p-value = 0.596