Middle Cerebral Artery Stenosis: Stenting Is One of the Options: No
Scott E. Kasner

Stroke. 2007;38:1420-1421; originally published online March 1, 2007;
doi: 10.1161/01.STR.0000259845.00988.2b
Stroke is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2007 American Heart Association, Inc. All rights reserved.
Print ISSN: 0039-2499. Online ISSN: 1524-4628

The online version of this article, along with updated information and services, is located on the
World Wide Web at:
http://stroke.ahajournals.org/content/38/4/1420

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in Stroke can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Stroke is online at:
http://stroke.ahajournals.org//subscriptions/
Middle Cerebral Artery Stenosis: Stenting Is One of the Options

No

Scott E. Kasner, MD

There is no doubt that stenting can widen a narrowed intracranial artery. However, serious questions remain about the role of this intervention in clinical practice. Medical technology often leaps into practice ahead of the science needed to support its widespread application. This is particularly common with new devices, which are not required to have demonstrated efficacy or safety with the same rigor as new medications. Moreover, they are not required to be superior to existing therapies. This is a flaw in the US regulatory system, and the premature approval of devices under humanitarian device exemptions often results in a “shoot first, ask questions later” approach that hampers subsequent attempts to determine whether these devices are actually beneficial.

The Warfarin versus Aspirin for Symptomatic Intracranial Disease (WASID) trial, the only large prospective study of intracranial stenosis, showed that patients with a stroke or transient ischemic attack attributable to a 50% stenosis of a major intracranial artery (internal carotid siphon, middle cerebral artery, vertebral artery, or basilar artery) confront a 12% per year risk of recurrent stroke in the territory of the stenosis, with the majority of strokes occurring in the first year. Despite longstanding beliefs that stenoses of some arteries pose different risks than others, WASID showed that patients with disease of the middle cerebral artery were no better or worse off than those with stenosis of another major artery. so I will address the treatment of all the major intracranial arteries based on the available evidence. Warfarin was not more effective than aspirin in prevention of recurrent stroke and carried a higher risk of serious bleeding and death. No subgroup of patients could be identified for whom warfarin was superior to aspirin. Prospectively, prespecified analysis demonstrated that patients with 70% stenosis or recent symptoms, particularly women, faced an enormous risk of recurrent stroke, possibly exceeding 20% in the first year. Some clinicians may interpret the WASID results nihilistically, and choose not to investigate the intracranial vessels because patients will just get aspirin as a default medication regardless of the findings. This approach is not advisable, because these patients probably require intensive risk factor modification at a minimum. Others with an eye toward intervention may feel compelled to offer these patients the most aggressive therapy possible, namely intracranial stenting.

The only device available in the US specifically for the treatment of intracranial atherosclerosis is the Wingspan stent made by Boston Scientific. The Wingspan self-expanding stent was studied in 45 patients with symptomatic intracranial (≥ 50%) stenosis who had recurrent cerebral ischemia on medical therapy. There were no medically treated control patients. The stent was successfully deployed in 44 of 45 patients and the periprocedural (30 day) risk of stroke or death was 4.4% (95% CI: 0.5% to 15%). Restenosis occurred at 6 months in 7.5%, and ipsilateral stroke or death at 1 year occurred in 9.3% (95% CI: 2.6% to 22%; Presented at the American Stroke Association International Stroke Conference, Kissimmee, Florida, February 2006). Other published small stenting registries have shown fairly similar results with devices not available or not approved in the US, but the total number of patients in these studies is only about 100 (including the 45 Wingspan patients). These data show that the arteries can be widened with reasonable safety, but do not show that this approach is better than medical therapy alone. Nevertheless, the FDA approved the Wingspan device under a humanitarian device exemptions. Unfortunately, the term “refractory” patients for whom stenting seems a logical option, must be viewed skeptically, because the patients probably require intensive risk factor modification at a minimum. Others with an eye toward intervention may feel compelled to offer these patients the most aggressive therapy possible, namely intracranial stenting.

Ultimately, the stenting registries and the WASID study populations may not be comparable, and it is still anyone’s guess whether one group was actually at greater baseline risk than the other.

Received December 10, 2006; accepted December 11, 2006.

From the Department of Neurology, University of Pennsylvania Medical Center, Philadelphia, Pa.

Correspondence to Scott E. Kasner, MD, Comprehensive Stroke Center, Department of Neurology, University of Pennsylvania Medical Center, 3400 Spruce St, Philadelphia, PA 19104. E-mail kasner@mail.med.upenn.edu

Stroke is available at http://www.strokeaha.org
DOI: 10.1161/01.STR.0000259845.00988.2b

© 2007 American Heart Association, Inc.

Stoke is available at http://stroke.ahajournals.org/ by guest on March 6, 2014
Stenting is an experimental procedure, though the FDA has created a loophole for clinical practice. This loophole should remain very small. At present, patients with intracranial stenosis should be treated with antiplatelet therapy. For the high-risk groups, there is great hope and hype about the efficacy of stenting. I share that hope, but a randomized trial of intracranial stenting (coupled with an intensive medical regimen for all) is urgently needed before enthusiasm becomes rampant for an unproven approach that merely fills a therapeutic void.

**Disclosures**

None.

**References**


Key Words: MCA  ■  stenting