

Ischaemic Stroke – Stenting versus Surgery for Carotid Disease

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Abstract

Extracranial internal carotid artery stenosis is one of the most common and best studied causes of stroke. Revascularisation with carotid endarterectomy (CEA) has been shown to be beneficial for patients with severe stenosis associated with stroke or transient ischaemic attack (TIA) and for many patients with moderate stenosis associated with stroke or TIA. CEA has also been shown to be beneficial for patients with asymptomatic severe stenosis if they have a reasonable expected lifespan and surgical risk, but the benefit is greater for men compared with women. Carotid angioplasty and stenting (CAS) has become a viable alternative procedure for carotid revascularisation with less risk of major bleeding complications and cranial nerve injury. Randomised studies of CEA versus CAS have found that the endovascular approach is associated with a lower risk of myocardial infarction but a higher risk of peri-procedural stroke which has a greater impact on long-term quality of life. Thus, recommending CEA or CAS must be based upon individual patient characteristics and their preferences, but at this point it appears that most patients should still be receiving CEA if an intervention is required.

Keywords

Stroke prevention, carotid stenosis, carotid endarterectomy, carotid angioplasty and stenting, carotid revascularisation

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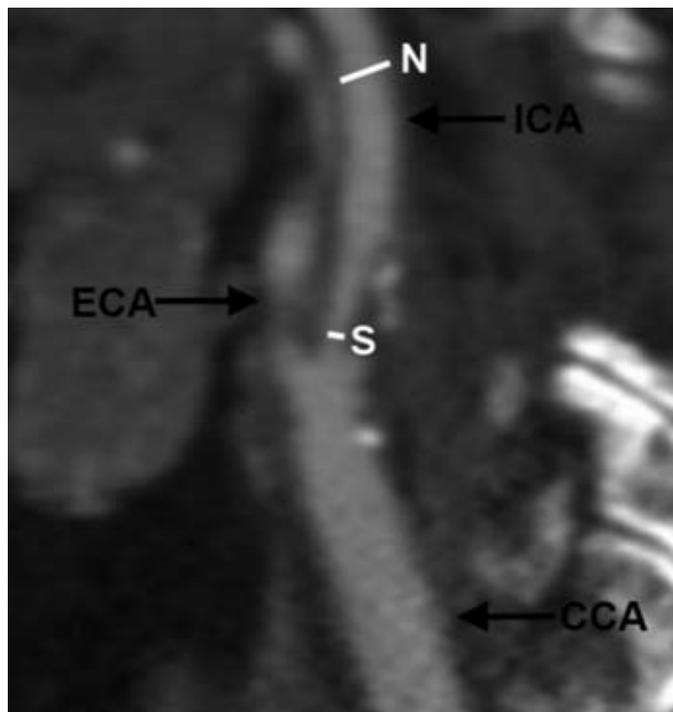
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The purpose of this article is twofold: first, to review the studies comparing carotid endarterectomy (CEA) with medical treatment to help decide who should undergo revascularisation; and secondly, to review studies comparing carotid angioplasty and stenting (CAS) versus CEA to see how they should be revascularised. Extracranial internal carotid artery stenosis is a leading cause of ischaemic strokes and transient ischaemic attacks (TIAs). It is estimated that extracranial atherosclerotic carotid disease is responsible for 15–20 % of strokes and treatments for extracranial internal carotid artery stenosis are among the best-studied interventions for preventing stroke.^{1–3} Several groundbreaking studies in the 1990s confirmed the benefit of surgical revascularisation for most patients with haemodynamically significant carotid stenosis. At this point, CEA is considered to be the gold standard treatment for symptomatic carotid stenosis and many patients with asymptomatic carotid stenosis also undergo revascularisation.^{4–10} With advances in endovascular techniques, CAS has evolved into a viable alternative to CEA and considerable interest has been shown in determining whether endovascular treatment is comparable to surgery for the treatment of carotid stenosis. Determining whether a carotid stenosis has been symptomatic or asymptomatic is essential to deciding whether an individual patient would benefit from a revascularisation procedure, as well as the urgency required to undertake such an intervention. Carotid artery stenosis is considered symptomatic if the patient has experienced focal neurological symptoms related to ischaemia in the ipsilateral retina causing monocular blindness, or in the ipsilateral cerebral hemisphere, potentially causing contralateral hemiparesis, hemianaesthesia, a visual field cut, and neglect in the non-dominant hemisphere, or aphasia in the dominant hemisphere.

Carotid Endarterectomy in Symptomatic Carotid Stenosis

In the 1990s, two large randomised controlled trials, namely the North American symptomatic carotid endarterectomy trial (NASCET)^{4,6} and the European carotid surgery trial (ECST),^{5,7} established that patients with symptomatic carotid stenosis benefit from CEA. NASCET was a randomised prospective multicentre trial carried out to assess the efficacy of CEA versus medical treatment in patients with symptomatic carotid atherosclerotic disease. The study enrolled 659 patients who had a hemispheric or retinal TIA or a non-disabling stroke within the 120 days before entry. The result showed a significant benefit of CEA in patients with 70–99 % symptomatic stenosis. The two-year ipsilateral stroke risk was 26 % in the medically treated patients versus 9 % in the surgical group ($p < 0.001$). The absolute risk reduction (ARR) was 17.0 % and the number needed to treat (NNT) was found to be six at two years. In patients with 50–69 % symptomatic stenosis, the benefit was more modest; the five-year rate of ipsilateral stroke was 15.7 % in patients treated with surgery and 22.2 % in patients who received medical treatment (ARR 6.5 %, NNT 15.4, $p = 0.045$). Finally, with < 50 % symptomatic stenosis, there was no significant difference, with a five-year rate of ipsilateral stroke of 14.9 % in the CEA group and 18.7 % in the medical therapy group ($p = 0.16$).^{4,6} Subset analysis found that patients who were aged 75 and older benefited more from CEA than younger patients.¹¹ *Post hoc* analyses further revealed gender differences in the 50–69 % group, with a statistical benefit from CEA seen only in men but not in women. ECST was a multicentre randomised controlled trial that enrolled 3,024 patients who had at least one transient or mild symptomatic ischaemic vascular event within the previous six months due to ipsilateral carotid artery stenosis.⁵ ECST initially used a different approach from NASCET

Figure 1: Measurement of an Internal Carotid Artery Stenosis Using the North American Symptomatic Carotid Endarterectomy Trial Criteria



The diameter of the artery in the projection that displays the greatest degree of stenosis (S) and at a normal segment distal to the stenosis (N). The percentage of stenosis = $(1-S/N) \times 100$. CCA = common carotid artery; ECA = external carotid artery; ICA = internal carotid artery.

to the measurement of the degree of carotid stenosis but they subsequently re-analysed the angiography data to be consistent with NASCET (see Figure 1 for NASCET criteria). Surgery reduced the five-year risk of any stroke or surgical death by 5.7 % (95 % confidence interval [CI] 0–11.6) in patients with 50–69 % stenosis by NASCET criteria (n=646, p=0.05) and by 21.2 % (95 % CI 12.9–29.4) in patients with 70–99 % stenosis by NASCET criteria without ‘near-occlusion’ (n=429, p<0.0001). Thus results of the ECST and NASCET were very consistent.¹²

Pooled Analysis and Subset Analysis of Carotid Endarterectomy Trials

Analysis of the pooled data from the NASCET, ECST, and the Veterans Affairs 309 study (a smaller randomised trial involving 189 patients with symptomatic carotid stenosis) confirmed the efficacy of CEA in patients with symptomatic carotid disease.¹³ The analysis showed that surgery increased the five-year risk of ipsilateral ischaemic stroke in patients with less than 30 % carotid stenosis, had no effect in patients with 30–49 % carotid stenosis, and was of marginal benefit in those with 50–69 % carotid stenosis. However, surgery was highly beneficial in patients with ≥70 % carotid stenosis but not near-occlusion. Importantly, surgical morbidity and mortality exceeding 6 % in symptomatic stenosis could negate the benefit gained from CEA.^{14,15} The combined data allowed for more precise subgroup analyses. For timing of the procedure, it was seen that CEA was most beneficial if carried out within the first two weeks after a non-disabling stroke or TIA.¹⁶ In general, men benefit from CEA more than women with symptomatic carotid stenosis revascularisation; however, CEA is clearly beneficial for women with 70–99 % symptomatic carotid stenosis.¹⁷ Some symptomatic subgroups appeared to derive more benefit from CEA and these include patients aged 75 years or more, patients with ulcerated plaques and patients with recent TIAs within two weeks of randomisation.¹⁶ CEA is also likely to be beneficial for patients

who have symptomatic ipsilateral carotid stenosis and co-existing severe contralateral carotid stenosis or occlusion, in spite of the increased risk compared with medical treatment alone.¹⁸ The interactions between all of these factors are complicated, but it is worth noting that time to CEA has the greatest impact on the potential benefit for women, such that it is reasonable to consider revascularising women with 50–69 % stenosis, if this is carried out within two weeks of the first ischaemic event.¹⁹ There was no clear benefit of the procedure in patients with total or near-total occlusion of the symptomatic ipsilateral internal carotid artery and it is unknown whether patients with ipsilateral stroke with disabling deficits or severe co-morbidities due to a medical or surgical condition would benefit or not.²⁰

Carotid Angioplasty and Stenting in Symptomatic Carotid Disease

Based upon the previously discussed studies, CEA is considered the gold standard therapy for patients with symptomatic severe carotid stenosis. CAS is an attractive alternative as it is less invasive and associated with less cranial nerve injury and fewer bleeding complications. Within the past five years, considerable high-level data have become available as multiple randomised studies that have compared CAS to CEA have been completed.

The Stent-protected angioplasty versus carotid endarterectomy (SPACE) trial was an international, multicentre, randomised controlled European study designed to test the non-inferiority of CAS to CEA for the treatment of severe symptomatic carotid stenosis. One thousand two hundred patients with symptomatic carotid artery stenosis were randomly assigned within 180 days of TIA or moderate stroke (modified Rankin scale score of ≤3) to carotid artery stenting (n=605) or CEA (n=595).²¹ The primary endpoint (rate of ipsilateral ischaemic stroke or death occurring within 30 days of the procedure) was 6.8 % in CAS and 6.3 % in CEA (absolute difference of 0.51 %, 90 % CI -1.89–2.91), which was greater than the predefined threshold. Thus, the study failed to prove the non-inferiority of CAS compared with CEA for the peri-procedural complication rate. In a *post hoc* analysis, older age in the CAS group (but not the CEA group) was significantly associated with an increased risk of ipsilateral stroke or death.²² After two years follow-up, there was no statistically significant difference between CAS and CEA in the composite endpoint of any peri-procedural stroke or death and ipsilateral ischaemic stroke up to two years after the procedure in both intention-to-treat (9.5 versus 8.8 %) and per-protocol (9.4 versus 7.8 %) analyses. The incidence of recurrent carotid stenosis ≥70 % at two years, as defined by ultrasound, was significantly higher after carotid artery stenting in both analyses (10.7 versus 4.6 % by intention-to-treat). However, it cannot be excluded that the degree of in-stent stenosis is slightly overestimated by conventional ultrasound criteria.²³

The Endarterectomy versus angioplasty in patients with symptomatic severe carotid stenosis (EVA-3S) was a French multicentre clinical trial that randomised 527 patients to endarterectomy (n=262) or CAS (n=265) to prove non-inferiority of CAS to CEA in low-risk patients with symptomatic carotid stenosis of ≥60 %. The 30-day incidence of any stroke or death, the composite primary outcome measure, was significantly higher with CAS than with CEA (9.6 versus 3.9 %, relative risk [RR] 2.5, 95 % CI 1.2–5.1) and the incidence of disabling stroke or death was 1.5 versus 3.4 % for the CEA and CAS groups, respectively. The trial was stopped prematurely due to an excess number of deaths in the CAS group.²⁴ At four years of follow-up, it was found that the cumulative probability of peri-procedural stroke or death and

non-procedural ipsilateral stroke was higher with stenting than with endarterectomy (11.1 versus 6.2 %, hazard ratio [HR] 1.97, 95 % CI 1.06–3.67, p=0.03). There were more major local complications after stenting and more systemic complications (mainly pulmonary) after endarterectomy, but the differences were not significant. Cranial nerve injury was more common after endarterectomy than after stenting.²⁴

The International carotid stenting study (ICSS) was a multicentre European trial, in which 1,713 patients (age >40 years) with recently symptomatic carotid artery stenosis were randomly assigned to receive carotid artery stenting (n=855) or CEA (n=858).²⁵ The 30-day risk of stroke, death, or myocardial infarction (MI) was significantly higher after CAS than after CEA (7.4 versus 4.0 %, RR 1.8, 95 % CI 1.2–2.8, p=0.003). The 120-day risk of stroke, death, or MI was still higher in the stenting cohort (8.5 versus 5.2 %; p=0.006). Moreover, in a subset of 231 patients in the ICSS who had brain magnetic resonance imaging (MRI), the proportion of patients with new ischaemic brain lesions on diffusion-weighted MRI at a median of one day after treatment was significantly higher in the stenting group than in the endarterectomy group (50 versus 17 %, odds ratio 5.2, 95 % CI 2.8–9.8, p<0.0001).²⁶

It is important to note that, relative to CEA, CAS is a less mature procedure and, as with any intervention, experience and improvements in techniques and devices have an impact on the potential risk and efficacy. Many proponents of CAS have argued that EVA-3S, SPACE and ICSS did not insure that the interventionists had adequate experience with CAS prior to enrolling patients in the study. That said, analyses of the experience of interventionists in these studies did not show a relationship with peri-procedural events.^{27–30}

Most recently, the Carotid revascularization endarterectomy versus stenting trial (CREST) was published.³¹ CREST was a North American randomized multicentre trial comparing CAS with CEA in both symptomatic and asymptomatic patients. The primary endpoint was the occurrence of stroke, death, or MI during the peri-procedural period and ipsilateral stroke up to four years. About half of the patients enrolled had an asymptomatic >60 % stenosis and half were symptomatic with >50 % stenosis. The CREST study attempted to address the issue of inexperienced interventionists by enforcing a credentialing lead-in period of up to 20 CAS procedures prior to enrolling patients in the study. Overall, there was no significant difference in the rates of the primary endpoint between CAS and CEA (7.2 versus 6.8 %, HR 1.11, 95 % CI 0.81–1.51, p<0.51) during long-term follow-up (median 2.5 years). An interaction with age and treatment was detected (p<0.02). Outcomes were slightly better after CAS for patients aged <70 years and better after CEA for patients aged >70 years. The proportion of patients developing stroke within 30 days of the procedure was significantly higher in the CAS than the CEA group (4.1 versus 2.3 %, HR 1.8, 95 % CI 1.1–2.8); on the other hand, the frequency of MI within 30 days of the procedure was significantly lower in the CAS group (1.1 versus 2.3 %, HR 0.5, 95 % CI 0.3–0.9). However, at one-year follow-up the quality of life was significantly diminished for patients who developed stroke compared with those with MI.³¹ For the subgroup of patients with symptomatic carotid disease, the peri-procedural rate of stroke and death was significantly higher for those assigned to stenting compared with endarterectomy (6.0 versus 3.2 %, HR 1.89, 95 % CI 1.1–3.2).³² In addition, CREST found that the difference in peri-procedural complications between CEA and CAS was accentuated in women. Peri-procedural events occurred in 35 (4.3 %) of 807 men assigned to carotid artery stenting compared with 40 (4.9 %) of 823 assigned to CEA (HR 0.90, 95 % CI 0.57–1.41) and 31 (6.8 %) of 455

Table 1: Symptomatic Carotid Disease – What Do the Guidelines Recommend?

Symptomatic Carotid Artery Stenosis	Level of Evidence
1. For patients with recent TIA or ischaemic stroke within the past six months and ipsilateral severe (70–99 %) carotid artery stenosis, CEA is recommended if the peri-operative morbidity and mortality risk is estimated to be <6 %	(Class I; level of evidence A)
2. For patients with recent TIA or ischaemic stroke and ipsilateral moderate (50–69 %) carotid stenosis, CEA is recommended depending on patient-specific factors, such as age, sex and co-morbidities, if the peri-operative morbidity and mortality risk is estimated to be <6 %	(Class I; level of evidence B)
3. When the degree of stenosis is <50 %, there is no indication for carotid revascularisation by either CEA or CAS	(Class III; level of evidence A)
4. When CEA is indicated for patients with TIA or stroke, surgery within two weeks is reasonable, rather than delaying surgery, if there are no contraindications to early revascularisation	(Class IIa; level of evidence B)
5. CAS is indicated as an alternative to CEA for symptomatic patients at average or low risk of complications associated with endovascular intervention when the diameter of the lumen of the internal carotid artery is reduced by >70 % by non-invasive imaging or >50 % by catheter angiography	(Class I; level of evidence B)
6. Among patients with symptomatic severe stenosis (>70 %) in whom the stenosis is difficult to access surgically, medical conditions are present that greatly increase the risk for surgery, or when other specific circumstances exist, such as radiation-induced stenosis or restenosis after CEA, CAS may be considered	(Class IIb; level of evidence B)
7. CAS in the above setting is reasonable when performed by operators with established peri-procedural morbidity and mortality rates of 4–6 %, similar to those observed in trials of CEA and CAS	(Class IIa; level of evidence B)
8. For patients with symptomatic extracranial carotid occlusion, EC/IC bypass surgery is not routinely recommended	(Class III; level of evidence A)
9. Optimal medical therapy, which should include antiplatelet therapy, statin therapy and risk factor modification, is recommended for all patients with carotid artery stenosis and a TIA or stroke as outlined elsewhere in this guideline	(Class I; level of evidence B)

The revised American Heart Association/American Stroke Association (AHA/ASA) guidelines published in 2011 for the prevention of stroke make the above recommendations for the management of symptomatic carotid stenosis. CAS = carotid angioplasty and stenting; CEA = carotid endarterectomy; EC/IC = extracranial/intracranial; TIA = transient ischaemic attack. Source: Furie et al., 2011.⁴¹

women assigned to carotid artery stenting compared with 16 (3.8 %) of 417 assigned to CEA (HR 1.84, 95 % CI 1.01–3.37, interaction p=0.064).³³

Asymptomatic Carotid Stenosis

Whether to recommend revascularisation for an asymptomatic carotid stenosis is a question that has persistently troubled neurologists, due to the fact that prior studies have found variable benefit from revascularisation, based upon a number of different patient characteristics. In addition, advances in medical therapies have raised doubts on whether the original CEA studies would show similar results if they were performed now.

Carotid Endarterectomy in Asymptomatic Carotid Stenosis

The Asymptomatic carotid atherosclerosis study (ACAS) was a prospective randomised multicentre trial that randomised 1,662 patients with asymptomatic carotid stenosis of >60 % to CEA and medical management.⁸ After a median follow-up of 2.7 years, the

Table 2: Asymptomatic Carotid Disease – What Do the Guidelines Recommend?

Asymptomatic Carotid Artery Stenosis	Level of Evidence
1. Patients with asymptomatic carotid artery stenosis should be screened for other treatable risk factors for stroke with institution of appropriate lifestyle changes and medical therapy	(Class I; level of evidence C)
2. Selection of asymptomatic patients for carotid revascularisation should be guided by an assessment of co-morbid conditions and life expectancy, as well as other individual factors and should include a thorough discussion of the risks and benefits of the procedure with an understanding of patient preferences	(Class I; level of evidence C)
3. The use of aspirin in conjunction with CEA is recommended, unless contraindicated, because aspirin was used in all of the cited trials of CEA as an antiplatelet drug	(Class I; level of evidence C)
4. Prophylactic CEA performed with <3 % morbidity and mortality can be useful in highly selected patients with an asymptomatic carotid stenosis (minimum 60 % by angiography, 70 % by validated Doppler ultrasound). It should be noted that the benefit of surgery may now be lower than anticipated based on randomised trial results and the cited 3 % threshold for complication rates may be high because of interim advances in medical therapy	(Class IIa; level of evidence A)
5. Prophylactic carotid artery stenting might be considered in highly selected patients with an asymptomatic carotid stenosis (>60 % on angiography, >70 % on validated Doppler ultrasonography, or >80 % on computed tomographic angiography or MRA if the stenosis on ultrasonography was 50–69 %). The advantage of revascularisation over current medical therapy alone is not well established	(Class IIb; level of evidence B)
6. The usefulness of CAS as an alternative to CEA in asymptomatic patients at high risk for the surgical procedure is uncertain	(Class IIb; level of evidence C)
7. Population screening for asymptomatic carotid artery stenosis is not recommended	(Class III; level of evidence B)

The revised American Heart Association/American Stroke Association (AHA/ASA) guidelines published in 2011 for the primary prevention of stroke make the above recommendations for the management of asymptomatic carotid disease. CAS = carotid angioplasty and stenting; CEA = carotid endarterectomy; MRA = magnetic resonance angiography. Source: Goldstein et al., 2011.⁴²

aggregate five-year risk of ipsilateral stroke, any peri-operative stroke, or death was estimated to be 5 versus 11 % for an RR reduction of 0.53 (95 % CI 0.22–0.72) favoring CEA. However, for surgery to be beneficial, the rate of peri-operative death and other serious complications had to be less than 3 %, and the expected patient survival had to be at least five years. The study showed higher incidence of peri-operative complications in women compared with men (3.6 versus 1.7 %), and men had an ARR of 8 % compared with 1.4 % in women, with women receiving no statistically significant benefit from revascularisation.

The Asymptomatic carotid surgery trial (ACST) was a subsequent randomised multicentre trial that enrolled 3,120 patients with ≥60 % asymptomatic carotid stenosis but no recent neurological symptoms (stroke or TIA) between immediate intervention (CEA) or indefinitely deferred CEA (until there was an associated stroke or TIA). Of 1,560 patients allocated to immediate treatment, half had CEA by one month and 88 % by one year and of the deferred group only 4 % per year underwent CEA.⁹ The CEA group had a peri-operative risk of stroke or death of 3.1 % within 30 days of surgery; however, the net five-year risk for all strokes or peri-operative death in the immediate CEA group was reduced by nearly half compared with the CEA deferral group (6.4 versus 11.8 %, 95 % CI 2.96–7.75), results that are similar to the ACAS study.

The ARR for preventing non-peri-operative stroke over five years was greater for men than for women (8.2 %, 95 % CI 5.64–10.78, versus 4.08 %, 95 % CI 0.74–7.41), although the benefit was statistically significant for both. CEA was shown to benefit patients <75 years of age, but there was no statistical benefit in patients who were older. ACST re-emphasised that, when selecting asymptomatic patients for carotid revascularisation, age, sex, life expectancy and the cited 3 % complication rate must all be taken into account. Finally, it is important to note that patients randomised to medical therapy in both ACAS and ACST were undertreated in terms of modern interventions such as statins and aggressive blood pressure goals. More recent prospective cohorts of medically treated patients with asymptomatic carotid stenosis have reported much lower stroke rates than these trials. A review of data collected from 11 studies showed that the average annual rates of ipsilateral or any other ischaemic stroke in asymptomatic severe carotid stenosis fell significantly in the last three decades with medical management alone.³⁴ It remains unknown whether modern aggressive medical management would be better than, or equivalent to, revascularisation in asymptomatic carotid stenosis, but it is an area which is gaining attention. Finally, it may be possible to identify patients at higher risk of stroke who then may be most likely to benefit from revascularisation. Based upon the ACST study, progression of carotid atherosclerosis on serial Doppler ultrasound studies is associated with increased risk of first stroke.³⁵ Furthermore, though not routinely available at every hospital, prolonged microembolic signal detection with transcranial Doppler (TCD) ultrasound has been validated in a multicentre study to accurately stratify high- and low-risk patients with asymptomatic carotid stenosis.³⁶

In summary, CEA can be recommended in men who have a life expectancy of at least five years with asymptomatic carotid stenosis of 60–99 %, provided the peri-operative risk of stroke and death is <3 %. In women, carotid revascularisation can be considered, particularly if they are younger and have a low expected peri-procedural risk.

Carotid Angioplasty and Stenting in Asymptomatic Carotid Stenosis – Stenting and Angioplasty with Protection in Patients at High Risk for Endarterectomy

Besides CREST, as discussed earlier, another study that looked into the utility of CAS in asymptomatic patients was the Stenting and angioplasty with protection in patients at high risk for endarterectomy (SAPPHIRE) trial. SAPPHIRE randomised 334 high-surgical-risk patients to receive CAS with embolic protection device or endarterectomy to test the hypothesis that CAS was not inferior to endarterectomy. These patients had symptomatic carotid stenosis of ≥50 % or asymptomatic carotid stenosis of ≥80 %; however, more than 70 % of patients had asymptomatic carotid disease.^{37,38} Patients were considered high risk for revascularisation based upon age >80, pulmonary or cardiac disease, prior neck surgery, or radiation. The primary endpoint of the cumulative incidence of a major cardiovascular event at one year, which included a composite of peri-procedural death, stroke, or MI (within 30 days after the procedure), and/or death or ipsilateral stroke between 31 days and one year, was 12.2 % for CAS compared with 20.1 % for CEA (absolute difference 7.9 %, 95 % CI -0.7–16.4 %). In the analysis of symptomatic patients with carotid stenosis (30 % in the CAS and 28 % in the endarterectomy group), the cumulative incidences of the primary endpoint at 30 days and also one year, in both groups were non-significantly different (2.2 versus 9.3 %, p=0.18 at 30 days and 16.8 versus 16.5 %, p=0.95 at one year). For patients with asymptomatic

carotid stenosis (70 % in the CAS and 72 % in the endarterectomy group), the cumulative incidence of the primary endpoint in the peri-procedural period was 5.4 versus 10.2 % (p=0.20) in the CAS and CEA groups, respectively. SAPHIRE hinted that CAS is not inferior to CEA in high-risk patients, namely with contralateral carotid occlusion, neck irradiation, prior neck surgery, severe cardiac/pulmonary disease, recurrent stenosis post-CEA and age more than 80 years.³⁹ In patients ≥80 years old, a meta-analysis of 41 studies of either CEA or CAS showed that the stroke rate was significantly higher for CAS compared with CEA (7.0 versus 1.9 %); the relative risks of death or MI at 30 days were fairly similar.⁴⁰ As mentioned above, half of the patients included in the CREST study had an asymptomatic carotid revascularisation. For these patients, the stroke and death rates were 2.5 ± 0.6 % for CAS and 1.4 ± 0.5 % for CEA (HR 1.88, 95 % CI 0.79–4.42, p=0.15).

Conclusions

The decision to recommend carotid revascularisation, by what method and in what timeframe, needs to be done on an individual basis depending on specific patient characteristics and the availability of surgeons and interventionists with a high volume of procedures and a track record with an acceptable complication rate. The accompanying tables provide the current recommendations from the American Heart Association regarding carotid revascularisation (see *Tables 1* and *2*). Clearly, the vast majority of patients with symptomatic carotid stenosis of 70–99 % will benefit from revascularisation, which should be carried out as soon as possible, assuming that the first event was not a major stroke. In this setting, it appears that CEA is more beneficial compared with CAS when the lesion is surgically accessible. CAS may be a

reasonable option in this setting if the lesion is surgically inaccessible, if there is restenosis post-CEA, radiation-induced stenosis, or underlying co-morbidities increasing surgical risk, assuming that the expected peri-procedural risk remains less than 6 %.

CEA does offer benefit, compared with medical treatment, for asymptomatic carotid stenosis ranging between 60 and 99 %, provided the life expectancy goal of five years is met and the combined peri-operative risk of stroke or death associated with the procedure is less than 3 %, though the benefit is greater in men compared with women. Based on the most recent data from CREST, CAS may be a reasonable choice in patients with an asymptomatic stenosis, particularly for younger men.

Lastly, it is important to bear in mind that the trials comparing medical therapy with CEA were designed and implemented over the last three decades. During this time, there have been significant changes in medical management for the prevention of stroke. With the advent of newer antiplatelet agents, more powerful statins with more aggressive lipid goals, use of non-beta-blocker blood pressure medications with more aggressive goals, decreased prevalence of smoking, and increased awareness of diet and exercise, there is ample evidence that the risk of stroke from an otherwise asymptomatic carotid stenosis is less than the risk seen in prior CEA studies. In some patients with asymptomatic stenosis, aggressive medical management alone rather than CEA could be offered with serial non-invasive imaging studies to ensure that the stenosis is not progressing. Microembolic signal detection with TCD may also help to risk-stratify asymptomatic patients. ■

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