Efficacy and safety of stenting for elderly patients with severe and symptomatic carotid artery stenosis: a critical meta-analysis of randomized controlled trials

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Objective: To investigate both short-term and long-term therapeutic efficacy and safety of carotid artery stenting (CAS) and carotid artery endarterectomy (CEA) for elderly patients with severe and symptomatic carotid artery stenosis.

Methods: PubMed, EMBASE, Cochrane Library, Clinical Trials Register Centers, and Google Scholar were comprehensively searched. After identifying relevant randomized controlled trials, methodological quality was assessed by using Cochrane tools of bias assessment. Meta-analysis was performed by RevMan software, and subgroup analyses according to different follow-up periods were also conducted.

Results: Sixteen articles of nine randomized controlled trials containing 6,984 patients were included. Compared with CEA, CAS was associated with high risks of stroke during periprocedural 30 days (risk ratio [RR]=1.47, 95% confidence interval [CI]: 1.15–1.88), 48 months (RR=1.37, 95% CI: 1.11–1.70), and >48 months (RR=1.76, 95% CI: 1.34–2.31). There was no significant difference in the aspects of death, disabling stroke, or death at any time between the groups. For other periprocedural complications, CAS decreased the risk of myocardial infarction (RR=0.44, 95% CI: 0.26–0.75), cranial nerve palsy (RR=0.09, 95% CI: 0.04–0.22) and hematoma (RR=0.31, 95% CI: 0.14–0.68) compared with CEA, while it increased the risk of bradycardia or hypotension (RR=8.45, 95% CI 2.91–24.58).

Conclusion: Compared with CEA, CAS reduced hematoma, periprocedural myocardial infarction, and cranial nerve palsy, while it was associated with higher risks of both short-term and long-term nondisabling stroke. And they seemed to be equivalent in other outcome measures. As regards to its minimal invasion, it should be applied only in specific patients.

Keywords: symptomatic carotid artery stenosis, carotid artery stenting, carotid artery endarterectomy

Introduction

According to the latest statistic from the American Heart Association, stroke ranks third among all the death causes, and every 4 minutes someone dies of stroke. Of all the strokes, 87% are ischemic, and people from 55 to 75 years of age who have a risk of stroke is 14% for men and 20% for women in the USA.¹

Carotid artery stenosis and occlusive diseases induced by many factors are important causes of ischemic stroke, and they often lead to immediate death although they count approximately 10%–15% of all the strokes.² Symptomatic patients with a >50% stenosis of vessel lumen was considered to be of high risk, and need to adopt aggressive treatments.³ Among the kinds of methods, carotid artery endarterectomy (CEA) was established as an effective option that periprocedural stroke/death...
is <6% for symptomatic and <3% for asymptomatic patients, and 10-year risk of stroke after CEA is approximately 2% per year.\textsuperscript{4,5} Carotid artery stenting (CAS) that emerged in the past 2 decades has also gradually developed to be an important and minimally invasive alternative. Compared with CEA, it was supposed to enhance recovery, reduce complication, and achieve cosmetic effect,\textsuperscript{4} and CAS was performed increasingly in clinical practice during the past few years.

However, therapeutic efficacy and safety of CAS compared with CEA were still uncertain. Up-to-date, a series of randomized controlled trials (RCTs) were designed and performed with various participants and follow-up periods.\textsuperscript{6–21} As insufficient statistical test power existed in single trials, the results and conclusions across the trials were controversial and confusing. Meanwhile, due to the lack of long-term results of follow-up longer than 2 years, current meta-analyses based on periprocedural and short-term data\textsuperscript{22,23} were not enough to provide valid and comprehensive evidence. Recently, in 2014 and 2015, many large-scale and multicenter RCTs stratified and published their long-term results ranged from postoperative 2–10 years.\textsuperscript{11,17,18}

It was necessary to take them together, and all the relevant RCTs involving short-term and long-term results could enhance our current knowledge and findings. So we conducted this critical and updated meta-analysis to conclude the comparative outcomes of CAS and CEA for carotid artery stenosis treatment.

Method

Literature search

A comprehensive search was performed on the databases including PubMed (1966.01–2015.05), EMBASE (1974.01–2015.05), and Cochrane Library (2015 Issue 5), as well as Clinical Trials Register Centers (up to 2015.05). Search terms were as follows: (“carotid artery” OR “vertebrobasilar” OR “cerebral” OR “cerebrovascular” OR “head and neck”) AND (“angiostenosis” OR “stenosis” OR “obstruct” OR “endothelial thicken” OR “occlusive disease”) AND (“stent” OR “stenting”). Medical subject headings were also used. Related articles, the references of relevant trials, and reviews were also screened to identify potential publications. Google Scholar was also searched for the last published articles.

Inclusion and exclusion criteria

Literature search results were first imported to citation manager software, and after duplication removed titles and abstracts were carefully scanned. At last, potential publications were further assessed by reading full-texts. Publications were included if 1) RCTs investigated the therapeutic efficacy and safety of CAS and CEA in carotid artery stenosis; 2) symptomatic patients >60 years, and with a severe carotid artery stenosis >50% of the luminal diameter, or asymptomatic patients with a >60% stenosis were participants; 3) preoperative aspirin was begun at least 72 hours before CAS or CEA and was continued indefinitely in both groups. Standard CEA was performed, and the stent used was self-expanding–nitinol stent with an emboli-protection device; and 4) primary outcomes should at least include stroke, death, or both of them. Secondary outcomes should include other complications such as transient ischemic attack, cranial nerve palsy, hematoma, restenosis, infection, and artery thrombosis. Meeting abstracts, reviews, non-RCTs, and non-English published papers were excluded.

Data collection and quality assessment

Reviewers extracted baseline characters of the included trials, which contained the first author, published year, case, average age, interventions, stenosis severity, diagnosis determination methods, and follow-up period. Data of outcomes were extracted in a pre-designed table for pooled analysis. Methodological quality was assessed by the tool of bias assessment provided by Cochrane Collaboration, which was based on six items:\textsuperscript{25} randomization, allocation concealment, participant, outcome assessment blinding, incomplete outcomes, selective reporting, and other bias. All data extraction and quality assessment were performed by two reviewers independently. Any disagreements were resolved by a third reviewer.

Statistical analysis

Meta-analysis was performed by using RevMan software (version 5.3, the Cochrane Collaboration, Copenhagen, Denmark). Subgroup analyzes were performed to identify important clinical characters, and all the analyzes were first performed based on clinical homogeneity. After that, $\chi^2$ and $F$ statistical tests were used to judge and present the statistical heterogeneity across the trials. A homogeneity was considered when $F \leq 50\%$, and fixed-effects model was chosen. Random-effects model was chosen when a heterogeneity existed, $F > 50\%$. Risk ratio (RR) and mean difference with their respective 95% confidence intervals (CIs) were presented for pooled effect size. Invested funnel plots were used to assess the risks of publication bias.

The meta-analysis was reported mainly according to Preferred Reporting Items for Systematic Reviews and
Meta-Analyses: the PRISMA statement.\textsuperscript{24} It did not involve any ethic issues.

**Results**

**Trial inclusion and quality assessment results**

Finally, 16 articles\textsuperscript{6–21} of nine trials containing 6,984 patients were included. There were 3,511 cases in the CAS group and 3,473 cases in the CEA group. Flow diagram of trial selection from initial search result to final decision is shown in Figure 1. The baseline characteristic of the included trials was presented in Table 1. Except three trials included both symptomatic and few asymptomatic patients,\textsuperscript{6,7,14,19,20} the others only included symptomatic patients. The three trials included patients who suffered >50% internal carotid artery (ICA) stenosis,\textsuperscript{6,7,18–20} and the four trials included patients who suffered >70% ICA stenosis.\textsuperscript{8,9,13,15–17} Follow-up ranged from postprocedural 30 days to 10 years.

Methodological quality assessment result was shown in Figure 2. The overall quality was good, whereas the item of blinding of participants and personnel was under unclear risk of bias. As a comparison of surgery, the procedure of CAS and CEA was really different, and blinding of participants and personnel was hard to realize.

**Primary periprocedural and follow-up results**

**Death**

According to the different follow-up period, a subgroup analysis including periprocedural 30 days, postprocedural 24, 48, and >48 months was conducted. Meta-analysis results

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**Table 1** Baseline characteristic of included randomized controlled trials

| Study          | Country    | Case (T/C, n) | Mean age (T/C, years) | Sex (M/F, n) | Intervention | Symptomatic Determination | Follow-up (T/C) | Sex (MF, n) | Mean age (T/C, years) |
|----------------|------------|---------------|-----------------------|--------------|--------------|--------------------------|----------------|-------------|---------------------|
| Gurm et al.\textsuperscript{6} | USA        | 167/167       | 72.5±8.3/72.6±8.9     | 111/54       | CAS/CEA      | >50% ICA stenosis        | Final 3 years  |            |                     |
| Yadav et al.\textsuperscript{7} | Switzerland| 10/10         | 69.8±6.7/71±5.9       | 8/2          | CAS/CEA      | >70% ICA stenosis        | Final 3 years  |      91/55    | 67.9±9.1±8.6±6.6   |
| Hoffmann et al.\textsuperscript{8} | Germany    | 43/44         | 67.9±9.1±8.6±6.6      | NR           | NR           | >60% ICA stenosis        | Final 3 years  |      122/15     | 67.6±8.7±6.6±6.6 |
| Steinbauer et al.\textsuperscript{9} | Germany    | 26/25         | 70.69                 | NR           | NR           | >70% ICA stenosis        | Final 3 years  |      188/74    | 67.7±9.1±8.6±6.6 |
| Mas et al.\textsuperscript{10–12} | France     | 80/80         | 66.8±9.2±8.7±6.7      | NR           | NR           | >70% ICA stenosis        | Final 3 years  |      206/59    | 68.7±9.1±8.6±6.6 |
| SPACE\textsuperscript{13} | Germany    | 65/65         | 70.69                 | 112/55       | CAS/CEA      | >70% ICA stenosis        | Final 3 years  |      206/59    | 68.7±9.1±8.6±6.6 |
| CA-VAST\textsuperscript{14} | Multicenter| 251/253       | 67.9±9.1±8.6±6.6      | NR           | NR           | >70% ICA stenosis        | Final 3 years  |      188/74    | 67.7±9.1±8.6±6.6 |
| ICSS\textsuperscript{15} | UK         | 53/53         | 70.66                 | NR           | CAS/CEA      | >70% ICA stenosis        | Final 3 years  |      174/77    | 67.7±9.1±8.6±6.6 |
| Brooks et al.\textsuperscript{16–17} | USA        | 83/83         | 70.69                 | NR           | CAS/CEA      | >50% CCA stenosis        | Final 3 years  |      174/77    | 67.7±9.1±8.6±6.6 |
| Silver et al.\textsuperscript{18} | USA and Canada | 1,262/1,240 | 68.9±9.0±8.7±8.6      | NR           | CAS/CEA      | >50% CCA stenosis        | Final 3 years  |      206/59    | 68.7±9.1±8.6±6.6 |

**Note:** Multicenter indicates countries all around the world.

**Abbreviations:** C, control group; CAS, carotid artery stenting; CEA, carotid artery endarterectomy; ICA, internal carotid artery; F, female; IQR, inter-quartile range; M, male; MRA, magnetic resonance angiogram; NR, not reported.
in fixed-effects model showed that there was no significant difference between CAS and CEA during periprocedural 30 days (F=0%, RR=1.22, 95% CI: 0.69–2.14, P=0.50), postprocedural 24 months (F=0%, RR=0.99, 95% CI: 0.71–1.37, P=0.93), 48 months (F=0%, RR=1.07, 95% CI: 0.89–1.29, P=0.48), and >48 months (F=0%, RR=1.23, 95% CI: 1.00–1.52, P=0.05), as shown in Figure 3.

**Stroke**

According to the different follow-up period, a subgroup analysis including periprocedural 30 days, postprocedural 24, 48, and >48 months was conducted. Meta-analysis results in fixed-effects model showed that CAS was associated with a higher stroke incidence during periprocedural 30 days (F=37%, RR=1.62, 95% CI: 1.31–2.00, P<0.0001), 48 months (F=0%, RR=1.37, 95% CI: 1.11–1.70, P=0.003), and >48 months (F=20%, RR=1.76, 95% CI: 1.34–2.31, P<0.0001) than CEA, whereas there was no significant difference between the groups during postprocedural 24 months (F=0%, RR=1.08, 95% CI: 0.80–1.47, P=0.60), as shown in Figure 4.

Subgroup analysis of the long-term effects also included periprocedural stroke incidence. To avoid a repeated analysis of periprocedural stroke incidence in postprocedural 24, 48, and >48 months, another subgroup analysis excluding periprocedural incidence was also conducted. It revealed that there was no significant difference between CAS and CEA during periprocedural 30 days to postprocedural 24 (F=0%, RR=0.98, 95% CI: 0.60–1.60, P=0.94) months and 48 months (F=0%, RR=1.07, 95% CI: 0.78–1.47, P=0.67). While, during periprocedural 30 days to postprocedural >48 months, the difference was statistically significant (F=44%, RR=1.58, 95% CI: 1.11–2.23, P=0.01).

**Myocardial infarction**

Subgroup analysis including periprocedural 30 days, postprocedural 12 and 36 months was performed. Meta-analysis results in fixed-effects model showed that compared with CEA, CAS achieved a decreased incidence of myocardial infarction (MI) during periprocedural 30 days (F=0%, RR=0.44, 95% CI: 0.26–0.75, P=0.003), whereas the difference between the groups did not reach statistical significance during postprocedural 12 months (F=0%, RR=0.41, 95% CI: 0.15–1.08, P=0.07) and 36 months (RR=0.64, 95% CI: 0.29–1.44, P=0.28), as shown in Figure 5.

**Disabling stroke and death**

Subgroup analysis including periprocedural 30 days, postprocedural 24, and >24 months was performed. Meta-analysis results in the fixed-effects model showed that there was no significant difference between the groups during periprocedural 30 days (F=0%, RR=1.19, 95% CI: 0.85–1.67, P=0.32), postprocedural 24 months (F=50%, RR=1.30, 95% CI: 0.93–1.82, P=0.13), and >24 months (F=48%, RR=1.02, 95% CI: 0.84–1.22, P=0.87), as shown in Figure 6.

**Other major complications**

Compared with CEA, CAS was associated with a significant decrease in periprocedural cranial nerve palsy (F=0%, RR=0.09, 95% CI: 0.04, 0.22, P<0.00001) and hematoma (F=41%, RR=0.31, 95% CI: 0.14, 0.68, P=0.003), whereas it was associated with a significant increase in bradycardia or hypotension (F=0%, RR=8.45, 95% CI: 2.91–24.58, P<0.0001). Besides, there was no significant difference in aspects of transient ischemic attack (F=11%, RR=1.58, 95% CI: 0.93–2.68, P=0.09), restenosis (F=0%, RR=2.22,
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95% CI: 0.51–9.60, \( P=0.29 \), arterial occlusion or thrombosis (\( F=23\% \), RR=1.49, 95% CI: 0.42–5.27, \( P=0.54 \)), and infection (\( F=0\% \), RR=0.60, 95% CI: 0.08, 4.54, \( P=0.62 \)), as shown in Figure 7.

Hospital stay

Three trials\(^{7,8,17}\) reported the data of hospital stay, and the meta-analysis in random-effects model showed that there was no significant difference between CAS and CEA (\( I^2=62\% \), mean difference \( -2.08 \), 95% CI: \( -4.47 \) to \( -0.32 \), \( P=0.09 \)).

Publication bias

Inverted funnel plots indicated that low risks of publication bias existed in the outcomes of death, stroke, and other major complications (Figure 8).

Discussion

The critical meta-analysis including 16 articles of nine RCTs with follow-up periods ranged from procedural 30 days to postprocedural 10 years. The pooled analysis altered that CAS was associated with increased risks of stroke compared with CEA during periprocedural 30 days and after postprocedural 4 years. And it confirmed the findings that higher risks of nondisabling stroke and bradycardia or hypotension, and a lower risk of MI were associated with CAS than CEA in the periprocedural period.

The estimated stroke rates were 6.19%, 9.79%, 9.56%, and 12.89%, respectively, at postprocedural 30 days, 2 years, 4 years and \( >4 \) years in the CAS group, compared with 3.82%, 9%, 6.97%, and 7.33% in the CEA group. Due to the loss to follow-up and the reduction of available cases, total stroke rate of both was increasing during follow-up period.
while CAS was always having a higher rate than CEA. In order to investigate the long-term effect of them, we further conducted a subgroup analysis including isolated data during procedural 30 days to final follow-up. The results revealed that CAS had a higher rate of stroke after postprocedural 4 years, based on the fact that there was no significant difference in outcomes of death and disabling stroke all the time. So it was clear that CAS had higher risks of nondisabling stroke during short-term of periprocedural 30 days, long-term of postprocedural 4 years, and overall period of follow-up, while a comparable risk during mid-term with CEA.

And CAS was demonstrated to achieve less cranial nerve palsy than CEA. However, it is hard for the patients in CEA group to identify a cranial nerve palsy from a stroke, so the actual stroke rate in CAS group might be even higher. Although the underlying mechanism was unclear, several studies reported that the increased incidence of stroke was mainly occurring in the contralateral carotid or vertebrobasilar territory. As is known, carotid artery and vertebrobasilar artery anastomoses each other through the circle of Willis (coW), and both the structure and function of coW are very important for blood supply of the whole brain. On the whole, local carotid artery surgeries may have different effects on the function of coW, and these different influences might be the major causes of therapeutic differences. A study including 139 patients reported that after carotid revascularization, the average diameter of ipsilateral precommunicating anterior cerebral artery (A1) increased 0.1 mm, while ipsilateral and contralateral posterior communicating artery (PCoA) decreased 0.12 mm and 0.08 mm. But, CAS led to much more diameter changes than CEA, with a maximum increase of 0.16 mm and decrease of 0.09 mm. After revascularization

| Study or subgroup                  | Experimental Events | Control Events | Weight | Risk ratio M–H, fixed, 95% CI | Risk ratio M–H, fixed, 95% CI |
|-----------------------------------|---------------------|----------------|--------|------------------------------|------------------------------|
| **Periprocedural 30 days**        |                     |                |        |                              |                              |
| Gurm et al*                       | 6                   | 5              | 167    | 3.9%                         | 1.20 (0.37, 3.86)             |
| Hoffmann et al*                   | 0                   | 1              | 10     | 1.2%                         | 0.33 (0.02, 7.32)             |
| EVA-3S**                         | 23                  | 21             | 259    | 5.4%                         | 3.26 (1.42, 7.47)             |
| SPACE**                          | 44                  | 37             | 589    | 29.1%                        | 1.15 (0.76, 1.76)             |
| CAVATAS**                        | 26                  | 22             | 253    | 17.0%                        | 1.19 (0.69, 2.04)             |
| ICSIS**                          | 59                  | 27             | 836    | 20.9%                        | 2.18 (1.40, 3.41)             |
| Brott et al*                      | 52                  | 29             | 1,240  | 22.6%                        | 1.76 (1.13, 2.76)             |
| **Subtotal (95% CI)**             | 3,395               | 3,354          | 100%   | 1.62 (1.31, 2.00)            |                              |
| Total events                      | 210                 | 128            |        |                              |                              |
| Heterogeneity: \(\chi^2=9.58, df=6, P=0.14\); \(I^2=37\%\) | | | | | |
| Test for overall effect: Z=4.43 \(P<0.00001\) | | | | | |
| **Postprocedural 24 months**      |                     |                |        |                              |                              |
| Gurm et al*                       | 15                  | 15             | 167    | 20.4%                        | 1.00 (0.51, 1.98)             |
| Steinbauer et al*                 | 1                   | 4              | 44     | 0.7%                         | 3.07 (0.13, 73.30)            |
| SPACE**                          | 64                  | 57             | 589    | 78.9%                        | 1.09 (0.78, 1.53)             |
| **Subtotal (95% CI)**             | 817                 | 800            | 100%   | 1.08 (0.80, 1.47)            |                              |
| Total events                      | 80                  | 72             |        |                              |                              |
| Heterogeneity: \(\chi^2=0.47, df=2, P=0.79\); \(I^2=0\%\) | | | | | |
| Test for overall effect: Z=0.53 \(P=0.60\) | | | | | |
| **Postprocedural 48 months**      |                     |                |        |                              |                              |
| Gurm et al*                       | 15                  | 15             | 167    | 11.1%                        | 1.00 (0.51, 1.98)             |
| EVA-3S**                         | 33                  | 17             | 262    | 12.7%                        | 1.92 (1.10, 3.36)             |
| CAVATAS**                        | 33                  | 27             | 253    | 20.0%                        | 1.23 (0.76, 1.99)             |
| Brott et al*                      | 105                 | 75             | 1,240  | 56.2%                        | 1.38 (1.03, 1.83)             |
| **Subtotal (95% CI)**             | 1,945               | 1,922          | 100%   | 1.37 (1.11, 1.70)            |                              |
| Total events                      | 186                 | 134            |        |                              |                              |
| Heterogeneity: \(\chi^2=2.40, df=3, P=0.49\); \(I^2=0\%\) | | | | | |
| Test for overall effect: Z=2.93 \(P=0.003\) | | | | | |
| **Postprocedural 48 months**      |                     |                |        |                              |                              |
| Steinbauer et al*                 | 4                   | 0              | 42     | 0.7%                         | 9.00 (0.50, 162.10)           |
| Brooks et al*                     | 4                   | 0              | 83     | 0.7%                         | 8.31 (0.45, 151.99)           |
| ICSIS**                          | 119                 | 72             | 857    | 98.6%                        | 1.66 (1.26, 2.19)             |
| **Subtotal (95% CI)**             | 985                 | 982            | 100%   | 1.76 (1.34, 2.31)            |                              |
| Total events                      | 127                 | 72             |        |                              |                              |
| Heterogeneity: \(\chi^2=2.49, df=2, P=0.29\); \(I^2=20\%\) | | | | | |
| Test for overall effect: Z=4.05 \(P<0.0001\) | | | | | |

Figure 4 Meta-analysis of periprocedural and postprocedural stroke.
Abbreviations: M–H, Mantel–Haenszel; CI, confidence interval.

Favors (experimental) Favors (control)
stenosis in carotid artery was eliminated, A1 perfusion was increased, and PCoA perfusion was back to normal. Although they were relatively small changes, they had marked hemodynamic effects. Still we cannot rule out the conclusion that the much more changes caused by CAS increased the higher risk of stroke than CEA because of the insufficient data and limitations from the study. While the other study demonstrated that the coW is not applicable. Test for overall effect: Z=1.07 (P=0.28)

| Study or subgroup | Experimental Events Total | Control Events Total | Weight | Risk ratio M–H, fixed, 95% CI |
|------------------|-------------------------|---------------------|--------|------------------------------|
| Periprocedural 30 days | | | | |
| Gurm et al | 4 | 167 | 10 | 167 | 22.9% | 0.40 (0.13, 1.25) |
| EVA-3S | 1 | 281 | 2 | 259 | 4.6% | 0.50 (0.06, 5.44) |
| CAVATAS | 0 | 251 | 3 | 253 | 8.0% | 0.14 (0.01, 2.77) |
| Brott et al | 14 | 1,262 | 28 | 1,240 | 64.6% | 0.49 (0.26, 0.93) |
| Subtotal (95% CI) | 1,941 | 1,919 | 100% | 0.44 (0.26, 0.75) |
| Total events | 19 | 43 | | |
| Heterogeneity: $\chi^2=0.70, df=3$ (P=0.67); $I^2=0\%$ |
| Test for overall effect: Z=3.01 (P=0.003) |

| Study or subgroup | Experimental Events Total | Control Events Total | Weight | Risk ratio M–H, fixed, 95% CI |
|------------------|-------------------------|---------------------|--------|------------------------------|
| Postprocedural 12 months | | | | |
| Gurm et al | 5 | 167 | 12 | 167 | 89.0% | 0.42 (0.15, 1.16) |
| Steinbauer et al | 0 | 43 | 1 | 44 | 11.0% | 0.34 (0.01, 8.14) |
| Subtotal (95% CI) | 210 | 211 | 100% | 0.41 (0.15, 1.08) |
| Total events | 5 | 13 | | |
| Heterogeneity: $\chi^2=0.01, df=1$ (P=0.91); $I^2=0\%$ |
| Test for overall effect: Z=1.81 (P=0.07) |

| Study or subgroup | Experimental Events Total | Control Events Total | Weight | Risk ratio M–H, fixed, 95% CI |
|------------------|-------------------------|---------------------|--------|------------------------------|
| Postprocedural 36 months | | | | |
| Gurm et al | 9 | 167 | 14 | 167 | 100% | 0.64 (0.29, 1.44) |
| Subtotal (95% CI) | 167 | 167 | 100% | 0.64 (0.29, 1.44) |
| Total events | 9 | 14 | | |

Figure 5 Meta-analysis of periprocedural and postprocedural myocardial infarction.

Abbreviations: M–H, Mantel–Haenszel; CI, confidence interval.

Figure 6 Meta-analysis of periprocedural disabling stroke and death.

Abbreviations: M–H, Mantel–Haenszel; CI, confidence interval.
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| Study or subgroup | Experimental Events Total | Control Events Total | Weight | Risk ratio M–H, fixed, 95% CI | Risk ratio M–H, fixed, 95% CI |
|-------------------|--------------------------|----------------------|--------|-----------------------------|-----------------------------|
| **Transient ischemic attack** | | | | | |
| Steinbauer et al\(^a\) | 3 | 43 | 2 | 44 | 9.6% | 1.53 (0.27, 8.74) |
| EVA-35\(^\text{t}\) | 6 | 261 | 2 | 159 | 12.1% | 1.83 (0.37, 8.95) |
| CAVATAS\(^\text{t}\) | 22 | 251 | 16 | 253 | 77.8% | 1.39 (0.75, 2.58) |
| Brooks et al\(^b\) | 1 | 53 | 0 | 511 | 0.5% | 28.44 (1.17, 689.70) |
| **Subtotal (95% CI)** | 608 | 967 | 100% | 1.58 (0.93, 2.68) | | |
| **Perioperative cranial nerve palsy** | | | | | |
| Gurn et al\(^a\) | 0 | 167 | 8 | 167 | 14.9% | 0.06 (0.00, 1.01) |
| Steinbauer et al\(^a\) | 0 | 43 | 1 | 44 | 2.6% | 0.34 (0.01, 8.14) |
| EVA-35\(^\text{t}\) | 3 | 261 | 20 | 259 | 35.2% | 0.15 (0.04, 0.49) |
| CAVATAS\(^\text{t}\) | 0 | 251 | 22 | 253 | 39.3% | 0.02 (0.00, 0.37) |
| Brooks et al\(^b\) | 0 | 53 | 4 | 51 | 8.0% | 0.11 (0.01, 1.94) |
| **Subtotal (95% CI)** | 775 | 774 | 100% | 0.09 (0.04, 0.22) | | |
| **Hematoma** | | | | | |
| Steinbauer et al\(^a\) | 1 | 43 | 6 | 44 | 22.9% | 0.17 (0.02, 1.36) |
| EVA-35\(^\text{t}\) | 1 | 261 | 2 | 259 | 7.8% | 0.50 (0.05, 5.44) |
| CAVATAS\(^\text{t}\) | 3 | 251 | 17 | 253 | 65.4% | 0.18 (0.05, 0.60) |
| Brooks et al\(^b\) | 3 | 53 | 1 | 51 | 3.9% | 2.89 (0.31, 26.85) |
| **Subtotal (95% CI)** | 608 | 607 | 100% | 0.31 (0.14, 0.68) | | |
| **Restenosis** | | | | | |
| Gurn et al\(^a\) | 2 | 167 | 1 | 167 | 40.1% | 2.00 (0.18, 21.85) |
| Hoffmann et al\(^b\) | 1 | 10 | 1 | 10 | 40.1% | 1.00 (0.07, 13.67) |
| Steinbauer et al\(^a\) | 2 | 43 | 0 | 44 | 19.8% | 5.11 (0.25, 103.51) |
| **Subtotal (95% CI)** | 220 | 221 | 100% | 2.22 (0.51, 9.60) | | |
| **Arterial occlusion or thrombosis** | | | | | |
| EVA-35\(^\text{t}\) | 4 | 261 | 1 | 259 | 25.1% | 3.97 (0.45, 35.27) |
| CAVATAS\(^\text{t}\) | 0 | 251 | 2 | 253 | 62.2% | 0.20 (0.01, 4.18) |
| Brooks et al\(^b\) | 1 | 53 | 0 | 51 | 12.7% | 2.89 (0.12, 69.32) |
| **Subtotal (95% CI)** | 565 | 563 | 100% | 1.49 (0.42, 5.27) | | |
| **Infection** | | | | | |
| Steinbauer et al\(^a\) | 0 | 43 | 1 | 44 | 59.6% | 0.34 (0.01, 8.14) |
| EVA-35\(^\text{t}\) | 1 | 261 | 1 | 259 | 40.4% | 0.99 (0.06, 15.78) |
| **Subtotal (95% CI)** | 304 | 303 | 100% | 0.60 (0.08, 4.54) | | |
| **Bradyarrhythmia or hypotension** | | | | | |
| EVA-35\(^\text{t}\) | 11 | 261 | 0 | 259 | 14.1% | 22.82 (1.35, 385.31) |
| Brooks et al\(^b\) | 19 | 53 | 3 | 51 | 85.9% | 6.09 (1.92, 19.36) |
| **Subtotal (95% CI)** | 314 | 310 | 100% | 8.45 (2.91, 24.58) | | |

**Figure 7** Meta-analysis of other perioperative complications.

**Abbreviations:** M–H, Mantel-Haenszel; CI, confidence interval.

plastic,\(^{23}\) except for inborn variation, stenting would alter the flow pattern, and nearly one-third of the subjects adopted CAS had a blockade of A1, PCoA or precommunicating posterior cerebral artery (P1) at postprocedural 1 week.\(^{23}\) Meanwhile, our meta-analysis mainly included patients of >50% ipsilateral stenosis, and most of them were symptomatic, who had high possibilities of variant structure and impaired function of coW compared with asymptomatic patients, although detailed
information is absent. Therefore, hemodynamic instability such as bradycardia or hypotension together with significantly altered coW flow pattern may to some extent explain the difference risk of nondisabling stroke between CAS and CEA.

Our meta-analysis also revealed that CAS had a significant high risk of stroke than CEA on long-term effects. In the current analysis, patients had an average age more than 67 years, and studies demonstrated that age was an independent predictor of stroke in CAS other than CEA. Patients >70 years old had a significantly increased risk of CAS in aspects of procedural stroke or death or ipsilateral stroke. After 48 months, patients including in the meta-analysis had an average age of nearly 71–72 years. So taking everything together, we may conclude that no matter to adopt or already adopted CAS, patients aged >70 years would suffer higher risk than CEA.

For other major complications, CAS significantly reduced the incidence of local hematoma. There was no significant difference in the aspects of infection, artery occlusive thrombosis, infection, restenosis, transient ischemic attack, and hospital stay. So as a minimally invasive surgery, CAS did not have obvious advantages than CEA, except for a decreased incision size, while clinically it seemed to be difficult for surgeons to perform a reoperation in recurrent patients who had undergone CAS.

Although the overall quality of the included RCTs was good, some other limitations existed: 1) stent material and type were not reported in detail, and different stent might have its special characters; 2) an accurate diagnosis of MI should be based on symptom, electrocardiogram, and Q-wave situation. However, most of the studies did not report a standard diagnosis method, and this might lead to potential bias; 3) surgeon’s experience may to affect the therapeutic effects. While, it was still without confirmed conclusions; 4) most of the patients were symptomatic, so the results and conclusions were mainly based on data of symptomatic patients. Their comparative efficacy on asymptomatic patients was inconclusive; 5) Stenosis location and coW structure may be the most important factors to influence future clinical judgment and choice, whereas currently RCTs did not involve them; and 6) only on one study performed cost analysis, while it only compared medical cost and did not add the cost of stent. Actually, CAS had a higher total cost than CEA.

**Conclusion**

CAS reduced hematoma, periprocedural MI, and cranial nerve palsy, while it was associated with a higher risk of nondisabling stroke of both short-term and long-term period in elderly patients with severe and symptomatic carotid stenosis. After considering age and survival time, we suggest that the choice on CAS or CEA in symptomatic patients should take into account coW situation, financial condition, and cosmetic requirement.

**Disclosure**

The authors report no conflicts of interest in this work.

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