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ESO Guideline on Endarterectomy and Stenting for Carotid Artery Stenosis

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Keywords:	carotid stenosis, endarterectomy, stenting, medical therapy, stroke, transient ischaemic attack
Abstract:	<p>Purpose: Atherosclerotic stenosis of the internal carotid artery is an important cause of stroke. The aim of this guideline is to analyse the evidence pertaining to medical, surgical and endovascular treatment of patients with carotid stenosis. These guidelines were developed based on the ESO standard operating procedure and followed the Grading of Recommendations, Assessment, Development, and Evaluation (GRADE) approach. The working group identified relevant questions, performed systematic reviews and meta-analyses of the literature, assessed the quality of the available evidence, and wrote recommendations.</p> <p>Findings: Based on moderate quality evidence, we recommend carotid endarterectomy (CEA) in patients with ≥ 60-99% asymptomatic carotid stenosis considered to be at increased risk of stroke on best medical treatment (BMT) alone. We also recommend CEA for patients with ≥ 70-99% symptomatic stenosis, and we suggest CEA for patients with 50-69% symptomatic stenosis. Based on high quality evidence, we recommend CEA should be performed early, ideally within two weeks of</p>

	<p>the last retinal or cerebral ischaemic event in patients with ≥ 50-99% symptomatic stenosis. Based on low quality evidence, carotid artery stenting (CAS) may be considered in patients < 70 years old with symptomatic ≥ 50-99% carotid stenosis.</p> <p>Discussion and conclusion: Several randomised trials supporting these recommendations were started decades ago, and BMT, CEA and CAS have evolved since. The results of another large trial comparing outcomes after CAS versus CEA in patients with asymptomatic stenosis are anticipated in the near future. Further trials are needed to reassess the benefits of carotid revascularisation in combination with modern BMT in subgroups of patients with carotid stenosis.</p>
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Declaration of conflicting interests

Name	Conflicts of interest
Leo H Bonati	Co-Principal Investigator of the ECST-2 and ACST-2 trial. Grants from the Swiss National Science Foundation, the Swiss Heart Foundation, the University of Basel and AstraZeneca for research related to carotid artery disease. Consultancy fees from AstraZeneca.
Joachim Berkefeld	Unrelated grants for angiographic imaging research from Siemens.
Gert Jan de Borst	Member of the writing committee of the ESVS guideline on the management of carotid atherosclerotic disease; EU Horizon 2020 grant for stratification of carotid disease; consultation fees from BAYER; Steering Committee Member ECST-2, CSTC.
Richard Bulbulia	Co-PI ACST-2, Co-Chair, ESC Position Paper on Carotid Disease
Hans-Henning Eckstein	Steering Committee SPACE-1, Co-PI SPACE-2, CSTC, Co-Investigator ROADSTER 1 and ROADSTER 2, Co-Investigator ACST-2, EU Horizon 2020 grant for stratification of carotid disease
Alison Halliday	PI for the ACST trials, Steering committee member for ECST-2, past TSC member ICSS. Chair-Elect of ESC Council on Stroke, co-author on the ESVS Carotid Guidelines, Co-author ESC Guidelines on Dyslipidaemias, CSTC
Isabelle van Herzele	Consulting and Research Grants from Silkroad Medical, Medtronic and Impact APV Europe.
Stavros Kakkos	Member of the writing committee of the ESVS guideline on the management of carotid atherosclerotic disease.
Igor Koncar	EU Horizon 2020 grant for stratification of carotid disease, Member of the writing committee of the ESVS guideline on the management of carotid atherosclerotic disease
Dominick J H McCabe	Prior grant funding from: The Meath Foundation; The Irish Institute of Clinical Neuroscience; The Irish Heart Foundation Stroke Prevention Bursary programme; The Trinity College Dublin Innovation Bursary; The Vascular Neurology Research Foundation, Ireland. Unrestricted educational grant funding from: Biogen Idec, Ireland; Verum Diagnostica, GmbH; Bayer HealthCare, Ireland; and SINNOWA Medical Science & Technology Co., China for unrelated translational research studies. Member of the writing committee for the ESVS guidelines for the management of patients with atherosclerotic carotid and vertebral artery stenosis (2017 and 2022).
Jean-Baptiste Ricco	Grant and consultation fees from BAYER, Member of the writing committee of the ESVS guideline on the management of carotid atherosclerotic disease
Peter Ringleb	Steering Committee Member SPACE-2, CSTC. Unrelated grant from Boehringer Ingelheim (ECASS-4). Unrelated lecture fees from Boehringer Ingelheim, Bayer, Pfizer, Daiichi Sankyo.

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Ethical Approval

Ethical approval was not necessary for the work described in this paper.

Informed consent

Not applicable.

Guarantor

A specific guarantor does not exist. The Module Working Group has jointly developed the manuscript.

Contributorship

All listed authors have contributed to the preparation and writing of the manuscript.

All authors researched literature and conceived the guideline. All authors were involved in protocol development. LB, SK and JB wrote the first draft of the manuscript. All authors reviewed and edited the manuscript and approved the final version of the manuscript.

ESO Guideline on Endarterectomy and Stenting for Carotid Artery Stenosis

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Abstract

Atherosclerotic stenosis of the internal carotid artery is an important cause of stroke. The aim of this guideline is to analyse the evidence pertaining to medical, surgical and endovascular treatment of patients with carotid stenosis. These guidelines were developed based on the ESO standard operating procedure and followed the Grading of Recommendations, Assessment, Development, and Evaluation (GRADE) approach. The working group identified relevant questions, performed systematic reviews and meta-analyses of the literature, assessed the quality of the available evidence, and wrote recommendations.

Based on moderate quality evidence, we recommend carotid endarterectomy (CEA) in patients with ≥ 60 -99% asymptomatic carotid stenosis considered to be at increased risk of stroke on best medical treatment (BMT) alone. We also recommend CEA for patients with ≥ 70 -99% symptomatic stenosis, and we suggest CEA for patients with 50-69% symptomatic stenosis. Based on high quality evidence, we recommend CEA should be performed early, ideally within two weeks of the last retinal or cerebral ischaemic event in patients with ≥ 50 -99% symptomatic stenosis. Based on low quality evidence, carotid artery stenting (CAS) may be considered in patients < 70 years old with symptomatic ≥ 50 -99% carotid stenosis.

Several randomised trials supporting these recommendations were started decades ago, and BMT, CEA and CAS have evolved since. The results of another large trial comparing outcomes after CAS versus CEA in patients with asymptomatic stenosis are anticipated in the near future. Further trials are needed to reassess the benefits of carotid revascularisation in combination with modern BMT in subgroups of patients with carotid stenosis.

Keywords:

carotid stenosis, endarterectomy, stenting, medical therapy, stroke, transient ischaemic attack

Introduction

Atherosclerotic carotid artery disease is one of the major causes of ischaemic stroke and transient ischaemic attack (TIA), accounting for about 10 to 15% of cases, depending on the method of aetiological classification and the patient population studied.(1) Atherosclerotic carotid stenosis mostly occurs at the carotid bifurcation, involving the distal common and the proximal internal carotid artery.(2) Other sites which are predisposed to develop atherosclerotic stenosis are the origin of the common carotid artery and the cavernous segment of the intracranial carotid artery. The prevalence of atherosclerotic carotid disease increases with age and is higher in men than in women. In Caucasian populations, $\geq 50\%$ stenosis of the carotid artery was identified in 2.3% of men in the 6th decade, in 6.0% in the 7th decade, and in 7.5% of men aged 80 years; in women, the corresponding prevalence figures were 2.0%, 3.6% and 5.0% in these age groups, respectively.(3)

This guideline provides recommendations on the use of carotid endarterectomy (CEA) and carotid artery stenting (CAS) in patients with symptomatic or asymptomatic stenosis of the extracranial carotid bifurcation caused by atherosclerosis. We did not review the available evidence regarding management of proximal common carotid artery or intracranial internal carotid artery stenosis, or non-atherosclerotic causes of stenosis, such as secondary to dissection, fibromuscular dysplasia, arteritis etc. Furthermore, we did not include aspects of diagnostic imaging, peri-procedural management, technical aspects of CEA and CAS, or medical therapy. Guidance on these topics can be found in other guidelines.(4-6)

Methods

This guideline document was commissioned by the European Stroke Organisation (ESO). A multi-disciplinary Module Working Group (MWG) was established, consisting of experts in the field from vascular neurology, vascular surgery and neuroradiology, who are represented as authors of this guideline document. The composition of this group was approved by the ESO Guidelines Board and the ESO Executive Committee, based on a review of the intellectual and financial disclosures of the proposed members.

The guidelines were developed using GRADE methodology(7) and the ESO Standard Operating Procedure.(8) In brief, we defined the patient population, the interventions and comparators, the outcomes of clinical interest (PICOs), and the design of studies to be included. The outcomes were rated as critical, important or of limited importance according to the GRADE criteria.(7, 8)

Population

This guideline makes recommendations on treatment of patients with symptomatic or asymptomatic atherosclerotic carotid stenosis. Carotid stenosis was defined as symptomatic if it had caused ischaemic cerebrovascular events in the ipsilateral eye (transient monocular blindness or retinal infarction) or cerebral hemisphere (transient ischaemic attack [TIA] or stroke) in the preceding 6 months. Asymptomatic carotid stenosis was defined as a stenosis which was not associated with any ocular or cerebral ischaemic events in the ipsilateral carotid territory within the preceding 6 months.

Patient subgroups

PICO questions were additionally analysed for following pre-specified patient subgroups when data were available:

1. Age ($< / \geq$ 70 years)
2. Sex
3. Degree of stenosis, according to the method used in the NASCET study(9) or its non-invasive equivalent (mild: $<50\%$, moderate: $50-69\%$, severe: $70-99\%$, near occlusion [defined as collapse of the distal lumen])
4. Time since most recent ischemic event (for symptomatic carotid stenosis)
5. Type of most recent ischemic event (for symptomatic carotid stenosis): stroke, transient ischemic attack, ocular ischaemia (including transient monocular blindness or *amaurosis fugax* and retinal infarction).

Interventions and Comparators

Interventions and comparators are CAS, CEA, and contemporary best medical therapy (as defined by the study authors at the time of the study). The guideline does not address carotid revascularisation done as part of acute stroke therapy, or carotid angioplasty without insertion of a stent.

Outcomes

We graded outcomes occurring in the peri-procedural period of carotid artery revascularisation, as well as outcomes occurring in the post-procedural period on a scale of 0-9 to classify them as either critical for decision making (grade 7-9; table 1); important, but not critical for making a decision (grade 4-6; table 1); or of limited importance for making a decision (grade 0-3). These grades were not included in the evidence profile.

The peri-procedural period was defined as the period between randomisation in the trial and 30 days after treatment, or as the first 30 days after randomisation in patients who did not undergo revascularisation (unless different definitions were used in individual trials in question). Peri-procedural outcomes were included as a measure of treatment safety. Post-procedural outcomes (i.e., outcomes occurring beyond the peri-procedural period) were included as a measure of treatment efficacy.

Peri-procedural and post-procedural outcomes graded as 'critical' or 'important' for decision making are listed in the table below and were included in the evidence profile.

Table 1: Outcomes

Peri-procedural outcomes graded as <i>critical</i> for decision making	<ul style="list-style-type: none"> - Death - Any stroke (ischaemic or haemorrhagic), defined as an acute onset of focal neurological function, with symptoms lasting for longer than 24 hours or leading to death within 24 hours, of non-traumatic vascular aetiology. Retinal infarction with visual loss lasting for longer than 24 hours, was included within the definition of stroke. - Major stroke, defined as resulting in substantial impairment or disability (measured by a modified Rankin scale(10) score of >2, typically 30 days or more after the event, if available), or death
Peri-procedural outcomes graded as <i>important</i> for decision making	<ul style="list-style-type: none"> - Myocardial infarction, according to the definitions used in the individual trials - Cranial nerve injury
Peri-procedural outcomes of limited importance	
Post-procedural outcomes graded as <i>critical</i> for decision making	<ul style="list-style-type: none"> - Ipsilateral stroke, occurring in the territory of the anterior or middle cerebral artery on the side of the randomised artery. - Any stroke - Major stroke, defined as resulting in substantial impairment or disability (measured by a modified Rankin Scale score(10) [mRS] of >2, if available), or death
Post-procedural outcomes graded as <i>important</i> for decision making	<p>Death</p> <p>Severe residual or recurrent stenosis (≥70% according to the NASCET method of grading stenosis (9) or its non-invasive equivalent) or occlusion of the treated artery.</p>

	Myocardial infarction, according to the definition used in the study
Post-procedural outcomes of limited importance	

Formation of PICO questions

A series of PICO (Population, Intervention, Comparator, Outcome) questions were developed and subsequently approved by the ESO Guidelines board and the ESO Executive Committee. The PICO questions were based on the peri-procedural and post-procedural outcomes, graded as critical or important for decision making, as well as combinations of these outcomes. We only compared peri-procedural outcomes on their own in trials of CAS versus CEA. This resulted in 4 PICO questions for the comparison of CEA versus medical therapy alone, 4 PICO questions for the comparison of CAS versus medical therapy alone, and 11 PICO questions for the comparison of CAS versus CEA in separate trials in patients with asymptomatic carotid stenosis and in patients with symptomatic carotid stenosis. We also formulated one PICO question concerning the risk of restenosis after CAS or CEA which was addressed using combined data from patients with symptomatic and asymptomatic carotid stenosis; these data are reported in the section on symptomatic carotid stenosis. Subgroup analyses for these PICO questions were also performed in the aforementioned pre-specified patient subgroups, where data were available.

Literature search, data extraction and synthesis

Literature searches were restricted to reports of randomised controlled trials (RCTs). We identified 3 systematic reviews of RCTs in the Cochrane Database of Systematic Reviews, which were of relevance to this guideline, one comparing CEA with medical therapy alone for asymptomatic carotid stenosis,(11) one comparing CEA with medical therapy alone for symptomatic carotid stenosis,(12) and one comparing CAS with CEA for asymptomatic or symptomatic carotid stenosis.(13) For the comparisons of CEA versus medical therapy, and CAS versus CEA, systematic searches of the MEDLINE, EMBASE, and Cochrane databases (from the date of the last search in the Cochrane reviews to August 10th 2020) were conducted by two ESO Guidelines methodologists (AL and MTR) using the same search terms which were defined in the Cochrane reviews. For the comparison of CAS versus best medical therapy, a *de novo* search of the literature was performed using the MEDLINE, EMBASE and Cochrane databases from their inception until August 10th 2020, using the search terms provided in the Appendix. To reduce the number of duplicate references identified, we simultaneously searched for relevant data in patients with asymptomatic and symptomatic carotid stenosis.

For each of the three main comparisons, a group of MWG members (a 'PICO group') was formed to select the studies for inclusion and to evaluate the available evidence. Within each PICO group, two MWG members independently screened the titles and abstracts of publications identified from the

searches (first level selection), and subsequently assessed the full text of potentially relevant studies (second level selection). Data were extracted independently by AL and MTR from studies which met criteria for second level selection, separately for patients with asymptomatic and those with symptomatic carotid stenosis. At least one additional MWG member checked the extracted data results for accuracy.

For some PICO questions (PICO 6.1 and 6.9), we included outcomes in pre-defined patient subgroups derived from pooled analyses of individual patient data (IPD) from the *EVA-3S*, *SPACE*, *ICSS* and *CREST* trials which were performed by the *Carotid Stenosis Trialists' Collaboration (CSTC)*.

The risks of selection, performance, detection, attrition and reporting bias in each randomised trial were assessed using the Cochrane Collaboration's tool.⁽¹⁴⁾ Heterogeneity across studies was assessed using Cochran's Q (reported as a p value) and I^2 statistics.⁽¹⁵⁾ For each PICO question and each outcome, the quality of evidence was rated using the GRADEpro Guideline Development Tool (McMaster University, 2015; developed by Evidence Prime, Inc.) as high, moderate, low or very low.⁽⁸⁾

The relevant PICO group was responsible for analysing the available data and formulating an evidence-based recommendation according to the GRADE evidence profiles and the ESO standard operating procedure. Random-effect meta-analyses were conducted and results were summarized as odds ratios (OR) and their 95% confidence interval. Heterogeneity across studies was assessed using the I^2 statistic. The absolute measure of intervention effects was calculated as the difference between the baseline risk of an outcome (patients receiving control intervention) and the risk of outcome after the intervention was applied (risk of an outcome in patients who received an intervention). Absolute effects are based on the relative magnitude of an effect with respect to the baseline risk, which is similar to risk differences. The *fewer value* represents any value below 1 per 1,000 and the *more value* represents any value more than 1 per 1,000.

The wording and the rating of the strength of each recommendation was passed by majority voting by all MWG members. An Expert Consensus Statement, based on voting by all MWG members, was presented where the PICO group considered that there was insufficient evidence available to provide clear evidence-based recommendations for situations in which practical guidance was needed for everyday clinical practice. Importantly, these Expert Consensus Statements should not be regarded as evidence-based recommendations since they only reflect the opinion of the majority of the members of the MWG.

The Guideline document was subsequently reviewed by all MWG members and modified until a consensus was reached. Finally, the guideline document was peer-reviewed and approved by external reviewers and members of the ESO Guidelines Board and ESO Executive Committee.

Unless specified otherwise, relative effects are given as risk ratios (RR) with 95% confidence intervals. Absolute effects are given as numbers of events per 1000 patients with 95% confidence intervals.

Results

1. Endarterectomy or medical therapy for asymptomatic carotid stenosis

Description of studies

The *Veterans Administration (VA) asymptomatic carotid stenosis cooperative study* randomised 444 men with $\geq 50\%$ asymptomatic carotid stenosis on angiography to CEA ($n=211$) or medical therapy alone ($n=233$) between 1983 and 1987.(16) Five per cent of patients turned out to have $<50\%$ stenosis after centralised analysis of the angiograms. Patients had never experienced any prior ipsilateral cerebrovascular events and were followed up for a mean of 47.9 months. The results were reported in 1993.

The *Asymptomatic Carotid Atherosclerosis Study (ACAS)* randomly allocated 1662 patients with $\geq 60\%$ asymptomatic carotid artery stenosis to CEA ($n=825$) or medical therapy alone ($n=834$) between 1987-1993. Patients were defined as being 'asymptomatic' if they never had cerebrovascular symptoms in the distribution of the "study" carotid artery or vertebrobasilar territory. Patients with contralateral cerebral hemispheric symptoms within the previous 45 days were excluded. Outcomes after a median follow-up period of 2.7 years were reported in 1995.(17) The definition of haemodynamically-significant carotid stenosis was based on meeting at least one of three pre-specified criteria from an ocular pneumoplethysmographic (OPG-Gee) examination, an ultrasound of carotid arteries and/or catheter angiography indicating a diameter stenosis of $\geq 60\%$ (NASCET methodology). Patients randomized to surgery on the basis of ultrasound findings, or ultrasound combined with OPG-Gee were also required to have a catheter angiogram prior to CEA. If a post-randomisation angiogram revealed that the contralateral carotid artery was more severely stenosed, that artery then became the allocated 'study artery'.

The *Asymptomatic Carotid Surgery Trial (ACST-1)* randomized 3120 patients with $\geq 60\%$ asymptomatic carotid stenosis on ultrasound to immediate CEA ($n=1560$, median delay 1 month [IQR 0.3–2.5]) or initial medical therapy with the option of deferred CEA ($n=1560$) between 1993 and 2003.(18, 19) The first ACST-1 report in 2004 provided data on outcomes during follow-up for up to 5 years (mean 3.4 years) after randomisation. (18) A subsequent report in 2010 included outcomes over a median follow-up period of 9 years (IQR 6–11 years) after randomisation.(19)

The *Aggressive Medical Treatment Evaluation for Asymptomatic Carotid Artery Stenosis (AMTEC)* study randomised 55 patients with 70-79% carotid stenosis to receive CEA ($n=31$) or medical therapy

alone (n=24) between 2009 and 2013.(20) Stenosis was graded by ultrasound examinations, but had to be confirmed by computed tomographic or magnetic resonance angiography (CTA/MRA) or catheter angiography. The trial was stopped prematurely by the independent data and safety monitoring board because of a high rate of the primary endpoint in the medical arm after a median follow-up period of 3.3 years (maximum, 5.0 years); results were reported in 2015.

Data from patients with 50-99% asymptomatic carotid stenosis randomly assigned to CEA (n=203) or medical therapy alone (n=113) between 2009 and 2013 in the three-arm *Stent-protected Angioplasty in Asymptomatic Carotid Artery Stenosis vs. Endarterectomy (SPACE-2)* trial were also included in the present section.(21, 22) A detailed description of the *SPACE-2* trial is provided in section 2 of the results.

The effects of treatment are presented with medical therapy as the reference group. A summary of findings is provided in table 2.

PICO 1.1: In patients with asymptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death?

There is moderate quality evidence that endarterectomy reduces the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death compared with medical therapy alone (RR 0.73; 95% CI: 0.59-0.90; equivalent to 19 fewer events with CEA per 1000, from 28 fewer to 7 fewer; figure 1.1).

PICO 1.2: In patients with asymptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of stroke in any territory, including peri-procedural death?

There is also moderate quality evidence that endarterectomy reduces the long-term risk of stroke in any territory, including peri-procedural death, compared with medical therapy alone (RR 0.74, 0.59-0.92; 31 fewer events with CEA per 1000 patients; from 48 fewer to 9 fewer; figure 1.2). Comparison of the data on the estimated number of ipsilateral strokes (PICO 1.1) and strokes in any territory (PICO 1.2) suggests that CEA might also prevent strokes occurring outside the territory supplied by the operated carotid artery.

Subgroup data regarding age, sex and severity of stenosis were derived from ACST-1 only. The effect of CEA is significantly modified by age (interaction p=0.04): there is moderate evidence of a benefit of

CEA in patients younger than 75 years (RR 0.62, 0.49-0.78; figure 1.2.1), but no evidence of benefit observed in patients ≥ 75 years old (RR 1.03, 95% CI 0.68 - 1.55, low quality evidence) (figure 1.2.2). There is no evidence of a modification of the effect of CEA according to sex (figure 1.2.1) or severity of stenosis (figure 1.2.3).

PICO 1.3: In patients with asymptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of major stroke, including peri-procedural death?

There is moderate quality evidence that endarterectomy reduces the long-term risk of major stroke, including peri-procedural death compared with medical therapy alone (RR 0.77: 0.61-0.98; 14 fewer events with CEA per 1,000; from 24 fewer to 1 fewer; figure 1.3).

PICO 1.4: In patients with asymptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of death?

There is no difference in long-term risk of death between patients assigned to endarterectomy and those assigned to medical therapy alone (RR 1.02, 95% CI: 0.88-1.20; 5 more events with CEA per 1000 patients, from 32 fewer to 53 more; low quality evidence; figure 1.4).

Analysis of current evidence and evidence-based recommendation

Data to assess the benefit of endarterectomy compared with medical therapy alone in patients with asymptomatic carotid stenosis were available from five RCTs which included a total of 5791 patients with mainly $\geq 60\%$ stenosis. We found moderate quality evidence that CEA reduces the risk of ipsilateral stroke and the risk of stroke in any territory in these patients. Based on the results of a single trial, we found no evidence that the benefit of CEA varied significantly between men and women, or according to the severity of the carotid stenosis. We did not find evidence of an increase of the benefit of surgery with increasing degree of asymptomatic carotid stenosis. However, a recent population-based study and systematic review suggested an increase in stroke risk with increasing degrees of asymptomatic carotid stenosis amongst patients receiving contemporary medical therapy.(23). Age influenced the effect of surgery in ACST-1, with benefits only observed in patients < 75 years of age. As the effect of age on treatment was only reported in a subgroup analysis of a single trial and taking into account the fact that cardiovascular disease mortality is decreasing and life expectancy is increasing in these patients, we refrained from making recommendations for CEA in patients with asymptomatic carotid stenosis based on fixed age limits.

The two largest trials contributing data were performed two to three decades ago. Best medical management of patients with atherosclerotic disease has evolved since, with more widespread use of statins and other lipid-lowering agents, and stricter control of blood pressure. Annual risks of ipsilateral stroke in more recent observational studies of patients with asymptomatic carotid stenosis range from 0.34-1.4%, which is lower than in the medical arms of the RCTs.(24-26) However, surgical techniques and peri-operative management have also improved since these landmark trials were completed. For these reasons, we downgraded the overall quality of evidence for indirectness.

Recommendation

In patients with $\geq 60\%$ asymptomatic carotid artery stenosis considered to be at increased risk of stroke on best medical therapy alone, we recommend carotid endarterectomy.

Quality of evidence: Moderate ⊕⊕⊕

Strength of recommendation: Strong for carotid endarterectomy ↑↑

This recommendation is independent of sex and stenosis severity.

Additional information

The question of whether carotid revascularisation confers additional benefits over modern medical therapy is being investigated in ongoing RCTs: the *Second European Carotid Surgery Trial* (ECST-2) enrolled 429 patients with asymptomatic or low-to-intermediate risk symptomatic carotid stenosis; follow-up is ongoing.(27) The *Carotid Revascularization and Medical Management for Asymptomatic Carotid Stenosis Trial* (CREST-2) includes two parallel trials of stenting vs. medical therapy and endarterectomy vs. medical therapy in patients with $\geq 70\%$ asymptomatic carotid stenosis(28).

There is debate about whether CEA should only be performed in patients with asymptomatic carotid stenosis who are considered to be at 'higher risk' of stroke on best medical treatment (BMT) alone. The guidelines published by the European Society for Vascular Surgery (ESVS) have proposed that surgery should be considered in selected patients with 60-99% asymptomatic carotid stenosis with one or more imaging or clinical characteristics that may be associated with an increased risk of late ipsilateral stroke.(4) These characteristics may include, among others, silent infarction on neuroimaging,(29) high degree(23) and progression of stenosis,(30, 31) echolucent plaque on ultrasound,(32, 33) intra-plaque haemorrhage on MRI,(34, 35) and micro-emboli(36) or reduced cerebrovascular reserve(37) on trans-cranial Doppler. This concept is currently being investigated in the *Endarterectomy Combined With Optimal Medical Therapy (OMT) vs OMT Alone in Patients With Asymptomatic Severe Atherosclerotic Carotid Artery Stenosis at Higher-than-average Risk of Ipsilateral*

Stroke (ACTRIS) trial, which is including patients with asymptomatic carotid stenosis who have imaging features believed to confer an increased risk of stroke.

Expert consensus statement

Expert consensus statement:


12/12 experts concluded that in selected patients 75 years of age or older with $\geq 60\%$ asymptomatic carotid artery stenosis and an expected survival of at least five years, who are considered to be at an increased risk of stroke on best medical therapy alone, carotid endarterectomy is suggested after careful consideration of the risks and benefits at a multi-disciplinary team meeting.

Table 2. Summary of findings for endarterectomy versus medical therapy for asymptomatic carotid stenosis (PICO 1.1-1.4).


Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		
PICO 1.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death												
5	randomised trials	not serious	not serious	serious ^a	not serious	none	139/2830 (4.9%)	190/2764 (6.9%)	RR 0.73 (0.59 to 0.90)	19 fewer per 1,000 (from 28 fewer to 7 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 1.2: Long-term risk of stroke in any territory, including peri-procedural death												
5	randomised trials	not serious	not serious	serious ^a	not serious	none	238/2830 (8.4%)	326/2764 (11.8%)	RR 0.74 (0.59 to 0.92)	31 fewer per 1,000 (from 48 fewer to 9 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 1.2.1a: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Men												
1	randomised trials	not serious	not serious	serious ^a	not serious	none	89/1021 (8.7%)	134/1023 (13.1%)	RR 0.67 (0.52 to 0.86)	43 fewer per 1,000 (from 63 fewer to 18 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 1.2.1b: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Women												
1	randomised trials	not serious	not serious	serious ^a	not serious	none	40/539 (7.4%)	65/537 (12.1%)	RR 0.61 (0.42 to 0.89)	47 fewer per 1,000 (from 70 fewer to 13 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 1.2.2a: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Age < 75 years												
1	randomised trials	not serious	not serious	serious ^a	not serious	none	98/1231 (8.0%)	160/1239 (12.9%)	RR 0.62 (0.49 to 0.78)	49 fewer per 1,000 (from 66 fewer to 28 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 1.2.2b: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Age < 75 years												
1	randomised trials	not serious	not serious	serious ^a	serious ^b	none	41/329 (12.5%)	39/321 (12.1%)	RR 1.03 (0.68 to 1.55)	4 more per 1,000 (from 39 fewer to 67 more)	⊕⊕○○ LOW	CRITICAL
PICO 1.2.3a: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Degree of stenosis < 80%												
1	randomised trials	not serious	not serious	serious ^a	not serious	none	56/641 (8.7%)	86/643 (13.4%)	RR 0.65 (0.48 to 0.90)	47 fewer per 1,000 (from 70 fewer to 13 fewer)	⊕⊕⊕○ MODERATE	CRITICAL

Certainty assessment							No of patients		Effect		Certainty	Importance
No of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		


PICO 1.2.3b: Long-term risk of stroke in any territory, including peri-procedural death. Subgroup: Degree of stenosis \geq 80% Stenosis

1	randomised trials	not serious	not serious	serious ^a	not serious	none	83/919 (9.0%)	113/917 (12.3%)	RR 0.73 (0.56 to 0.96)	33 fewer per 1,000 (from 54 fewer to 5 fewer)	 MODERATE	CRITICAL
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PICO 1.3: Long-term major stroke, including peri-procedural death

4	randomised trials	not serious	not serious	serious ^a	not serious	none	119/2619 (4.5%)	153/2531 (6.0%)	RR 0.77 (0.61 to 0.98)	14 fewer per 1,000 (from 24 fewer to 1 fewer)	 MODERATE	CRITICAL
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PICO 1.5: Long-term risk of death

4	randomised trials	not serious	not serious	serious ^a	serious ^b	none	699/2619 (26.7%)	667/2531 (26.4%)	RR 1.02 (0.88 to 1.20)	5 more per 1,000 (from 32 fewer to 53 more)	 LOW	IMPORTANT
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CI: Confidence interval; RR: Risk ratio

Explanations

a. Endarterectomy and medical therapy have evolved since the trials contributing the evidence were performed

b. Few events and wide confidence intervals

2. Stenting or medical therapy for asymptomatic carotid stenosis

Description of studies

The *Stent-protected Angioplasty in Asymptomatic Carotid Artery Stenosis vs. Endarterectomy (SPACE-2)* trial was a randomised multi-centre study in Germany, Austria and Switzerland which aimed to assess the safety and efficacy of CAS or CEA compared with best medical therapy (BMT) alone in patients with asymptomatic $\geq 50\%$ common or internal carotid artery stenosis. (22) Stenoses were considered asymptomatic if patients had not experienced ipsilateral amaurosis fugax, a TIA or stroke within the preceding 180 days. SPACE-2 started in 2009 as a three-arm trial randomly assigning patients to CEA+BMT, CAS+BMT, or BMT alone in a 3:3:1 ratio, with a target sample size of 3550 patients. For CAS, the use of protection devices was not mandatory. The trial design was changed in 2013 to a two-arm trial of CEA+BMT versus CAS+BMT. Due to slow recruitment, the trial was stopped prematurely in 2014 after 513 patients had been randomised to CEA (n = 203), CAS (n = 197), or BMT (n = 113). This section of the guidelines only includes outcomes of patients in the CAS and BMT groups. Results after 1 year of follow-up were previously published. The primary efficacy endpoint (the cumulative risk of any stroke or death from any cause within 30 days, plus any ipsilateral ischaemic stroke within five years of follow-up) is yet to be reported.

We excluded two smaller RCTs because these studies did not report outcomes by symptom status,(38, 39) or patients were treated with primary balloon angioplasty.(39) Therefore, the *SPACE-2* data were the only data which could be used to address the PICO questions in this section.

The effects of treatment are presented with medical therapy as the reference group. A summary of findings is provided in table 3.

PICO 2.1: In patients with asymptomatic carotid stenosis, does stenting compared with medical therapy alone reduce the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death?

There is very low quality of evidence from *SPACE-2* of a non-significant increase in the risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death with stenting compared with medical therapy alone (RR 3.44, 95% CI: 0.42-28.23; equivalent to 22 more events with CAS per 1000 patients, from 5 fewer to 241 more; figure 2.1).

PICO 2.2: In patients with asymptomatic carotid stenosis, does stenting compared with medical therapy alone reduce the long-term risk of stroke in any territory, including peri-procedural death?

There is also very low quality evidence from SPACE-2 of a non-significantly higher risk of stroke in any territory, including peri-procedural death with stenting compared with medical therapy (RR 4.59, 0.58-36.22; 32 more events with CAS per 1,000 patients, from 4 fewer to 312 more; figure 2.2).

PICO 2.3: In patients with asymptomatic carotid stenosis, does stenting compared with medical therapy alone reduce the long-term risk of major stroke, including peri-procedural death?

Only one such composite event occurred in each of the stenting and medical therapy groups in SPACE-2 (RR 0.57, 0.04-9.08; low quality evidence; figure 2.3).

PICO 2.4: In patients with asymptomatic carotid stenosis, does stenting compared with medical therapy alone reduce the long-term risk of death?

There is very low quality of evidence that the long-term risk of death did not differ between patients treated with stenting and medical therapy in SPACE-2 (RR 0.29, 0.05-1.54; figure 2.4).

Analysis of current evidence and evidence-based recommendations

The evidence from this single, prematurely terminated RCT is very limited. The recruited study population is too small, and the available follow-up period is too short to reliably compare data between treatment groups. We downgraded the evidence for the risk of bias (due to the early termination), imprecision, and indirectness (insufficient length of follow-up), resulting in a very low quality of evidence.

Recommendation

In patients with asymptomatic carotid stenosis, we recommend against carotid artery stenting as a routine alternative to best medical therapy alone.

Quality of evidence: Very low ⊕

Strength of recommendation: Weak against carotid stenting ↓

Recommendations regarding the choice between stenting and endarterectomy in patients with asymptomatic carotid stenosis, in whom revascularisation is considered to be appropriate are provided in chapter 3.

Additional information

Carotid artery stenting versus best medical therapy alone are being compared in one of the two parallel study arms in the ongoing *Carotid Revascularization and Medical Management for Asymptomatic Carotid Stenosis Trial (CREST-2)* (28).

Table 3. Summary of findings for stenting versus medical therapy for asymptomatic carotid stenosis (PICO 2.1-2.5).

Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Stenting	Medical therapy	Relative (95% CI)	Absolute (95% CI)		
PICO 2.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death												
1	randomised trials	serious ^a	not serious	serious ^b	very serious ^c	none	6/197 (3.0%)	1/113 (0.9%)	RR 3.44 (0.42 to 28.23)	22 more per 1,000 (from 5 fewer to 241 more)	⊕○○○○ VERY LOW	CRITICAL
PICO 2.2: Long-term risk of stroke in any territory, including peri-procedural death												
1	randomised trials	serious ^a	not serious	serious ^b	very serious ^c	none	8/197 (4.1%)	1/113 (0.9%)	RR 4.59 (0.58 to 36.22)	32 more per 1,000 (from 4 fewer to 312 more)	⊕○○○○ VERY LOW	CRITICAL
PICO 2.3: Long-term risk of major stroke, including peri-procedural death												
1	randomised trials	serious ^a	not serious	serious ^b	not serious ^c	none	1/197 (0.5%)	1/113 (0.9%)	RR 0.57 (0.04 to 9.08)	4 fewer per 1,000 (from 8 fewer to 72 more)	⊕⊕○○○ LOW	CRITICAL
PICO 2.5: long-term risk of death												
1	randomised trials	serious ^a	not serious	serious ^b	very serious ^c	none	2/197 (1.0%)	4/113 (3.5%)	RR 0.29 (0.05 to 1.54)	25 fewer per 1,000 (from 34 fewer to 19 more)	⊕○○○○ VERY LOW	IMPORTANT

CI: Confidence interval; RR: Risk ratio

Explanations

a. Trial was stopped early

b. Insufficient length of follow-up to assess long-term effects

c. Very wide confidence intervals

3. Stenting or endarterectomy for asymptomatic carotid stenosis

Description of studies

A single-centre trial in Lexington, Kentucky, USA randomised 85 participants with $\geq 80\%$ asymptomatic carotid stenosis to receive either CAS without a cerebral protection device (CPD) or CEA and reported results up to four years after randomisation in 2004.(40) . A further report in 2014 combined long-term outcomes for up to 10 years in both asymptomatic and symptomatic patients who were enrolled in another trial at the same institution, but the authors did not present separate data according to symptom status. (41) Therefore, we chose the 2004 report to extract outcome data from patients with asymptomatic stenosis to address our PICO questions.

The *Carotid Revascularization Endarterectomy versus Stenting Trial (CREST)*, a multicentre trial in the USA and Canada, randomised 1321 patients with $\geq 50\%$ symptomatic carotid stenosis and 1181 patients with $\geq 60\%$ asymptomatic carotid stenosis to CAS or CEA between 2000-2008.(42-49) Interventionists with an experience of < 30 CAS procedures were required to complete a training programme. The use of a CPD was mandatory during stenting. Initial results were published in 2010; the final trial results were published in 2016 with follow-up data for up to 10 years after randomisation (median of 7.4 years). Only data from asymptomatic patients were extracted for our analyses to address these PICO questions.

A single-centre trial in Houston, Texas, USA randomised 60 patients with $\geq 80\%$ asymptomatic carotid stenosis to receive CAS (with mandatory use of a CPD) or CEA. The primary outcome was 'cognitive performance' after treatment; this and other clinical outcome data for up to 6 months after randomisation were reported in 2014.(50) No data were available for 5 patients who withdrew consent or were lost to follow-up.

A single-centre trial conducted in Ostrava, Czech Republic, randomised 63 patients with asymptomatic and 87 patients with symptomatic $\geq 70\%$ carotid stenosis to undergo CAS (with the use of a CPD, where possible) or CEA and reported results in 2014.(51) The primary outcome was the occurrence of new ischaemic brain lesions on magnetic resonance imaging after treatment. Clinical outcome events up to 30 days after treatment were also reported, and these were made available and categorised according to symptom status following correspondence with the investigators.

The *Randomized Trial of Stent versus Surgery for Asymptomatic Carotid Stenosis (ACT-1)* allocated 1453 participants < 80 years of age with $\geq 70\%$ asymptomatic carotid stenosis in a 3:1 ratio to undergo

CAS (with mandatory use of a CPD) or CEA between 2005 and 2013.(52) A prior experience of ≥ 25 procedures was required from surgeons and interventionists. The initially planned sample size was 1658 participants, but the study was stopped prematurely due to slow enrolment. Results up to 5 years after randomisation were previously published.

A single-centre trial at *Carmel Medical Center* in Israel randomised 136 participants with $\geq 70\%$ asymptomatic carotid stenosis to receive CAS (with mandatory use of a CPD) or CEA. Results up to 5 years after randomisation were reported in 2017.(53) Three patients were lost to follow-up.

Events occurring up to 1 year after treatment were also extracted from the CAS and CEA groups of the 3-arm *SPACE-2* trial (described in results section 2).(22)

We did not include data from the multi-centre *Stenting and Angioplasty with Protection in Patients at High Risk for Endarterectomy (SAPPHIRE)* trial conducted in the USA,(54-56) from one Chinese multi-centre trial,(57) and two single-centre studies conducted in Beijing, China.(58, 59) Reasons for exclusion of these randomised studies were the inclusion of patients with both asymptomatic and symptomatic carotid stenosis without reporting of separate outcome data according to symptomatic status, inclusion of 'high surgical risk' patients only, or results in the English language only being available as a conference abstract .

The effects of treatment are presented with endarterectomy as the reference group. A summary of findings is provided in table 4.

PICO 3.1: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death?

There is moderate quality evidence that stenting is likely associated with an increased long-term risk of post-procedural ipsilateral stroke, peri-procedural stroke in any territory, or peri-procedural death (RR 1.25, 95%CI 0.88-1.79; equivalent to 9 more events with CAS per 1000 patients, from 4 fewer to 28 more; figure 3.1).

PICO 3.2: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of post-procedural ipsilateral stroke?

There is low quality evidence that endarterectomy and stenting do not differ in preventing post-procedural ipsilateral stroke, excluding peri-operative events (RR 1.12, 0.62-2.00; 3 more events with stenting per 1000 patients, from 8 fewer to 22 more; figure 3.2).

PICO 3.3: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of stroke in any territory, including peri-procedural death?

There is moderate quality evidence that stenting is likely associated with an increased long-term risk of stroke in any territory or peri-procedural death (RR 1.22, 0.87-1.71; 13 more events with stenting per 1000 patients, from 8 fewer to 42 more; figure 3.3).

PICO 3.4: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of major stroke, including peri-procedural death?

There is low quality evidence that endarterectomy and stenting do not differ in the long-term risk of major stroke or peri-procedural death (RR 0.99, 0.15-6.68; 0 fewer events with stenting per 1000 patients, from 20 fewer to 20 more; figure 3.4).

PICO 3.5: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of death?

There is low quality evidence that endarterectomy and stenting do not differ in the long-term risk of death (RR 0.82, 0.31-2.20; 5 fewer events with stenting per 1000 patients, from 18 fewer to 32 more; figure 3.5).

PICO 3.6: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural stroke?

There is moderate quality evidence that stenting is likely associated with an increased risk of peri-procedural stroke (RR 1.70, 0.99-2.93; 10 more events with stenting per 1000 patients, from 0 fewer to 28 more; figure 3.6).

PICO 3.7: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural death?

There is high quality evidence that endarterectomy and stenting do not differ in the risk of peri-procedural death (RR 0.33, 0.02-5.33; 1 less event with stenting per 1000 patients, from 1 less to 6
28

more; figure 3.7). We did not downgrade the quality of evidence for imprecision because only a single event occurred in each treatment group.

PICO 3.8: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural stroke or death?

There is moderate quality evidence that stenting is likely associated with an increased risk of peri-procedural stroke or death as compared to endarterectomy (RR 1.62, 0.96-2.76; 9 more events per 1000 patients, from 1 less to 27 more; figure 3.8).

PICO 3.9: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural major stroke or death?

There is moderate quality evidence that stenting is likely associated with a slight increase of the risk of major peri-procedural stroke or death (RR 1.54, 0.39-6.07; 2 more events with stenting per 1000 patients, from 2 fewer to 15 more; figure 3.9).

PICO 3.10: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural myocardial infarction?

There is low quality evidence that stenting is likely associated with a lower risk of peri-procedural myocardial infarction as compared to endarterectomy (RR 0.53, 0.25-1.15; 6 fewer events with stenting per 1000 patients, from 9 fewer to 2 more; figure 3.10). We additionally downgraded the quality of evidence for indirectness because all extracted events originated from the CREST and ACT-1 trials, where screening with ECG and cardiac enzymes of all patients was performed before and after treatment; the definition of myocardial infarction included elevation of cardiac enzymes alone, or in combination with ECG changes only (without clinical symptoms).

PICO 3.11: In patients with asymptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural cranial nerve injury?

There is high quality evidence that stenting is associated with a lower risk of peri-procedural cranial nerve injury than endarterectomy (RR 0.09, 95% CI 0.03-0.28; 30 fewer events per 1000 patients with stenting, from 32 fewer to 24 fewer; figure 3.11). We upgraded the quality of evidence by 2 levels for strength of effect.

Analysis of current evidence and evidence-based recommendation

Data comparing the short-term risks and long-term effects between stenting and endarterectomy for asymptomatic carotid stenosis were available from 7 trials including a total of 3373 patients. Most studies required patients to have $\geq 60\%$ carotid stenosis for inclusion. Duration of follow-up in the largest trials was for 5 years or more. The risks of most outcome events were low, which led us to downgrade the level of evidence for imprecision. Low event rates also precluded meaningful subgroup analyses. Overall, we found no clear evidence of statistically significant differences in outcomes between endarterectomy or stenting that were rated as critical for decision making when treating patients with asymptomatic carotid stenosis (low to moderate quality evidence). As the available evidence is not sufficient to recommend stenting as an alternative to endarterectomy, carotid endarterectomy presently remains the treatment of choice for patients with asymptomatic carotid stenosis considered to require revascularisation.

Recommendation

In patients with asymptomatic carotid stenosis in whom revascularisation is considered to be appropriate, we suggest endarterectomy as the current treatment of choice.

Quality of evidence: Moderate ⊕⊕⊕

Strength of recommendation: Weak for carotid endarterectomy ↑

Additional information

The Asymptomatic Carotid Surgery Trial-2 (ACST-2) has recently completed recruitment of 3.638 patients with asymptomatic carotid stenosis who were randomly assigned to CAS or CEA.(60) First results are expected in late 2021 and will considerably increase the evidence base, which may lead to updates to the above recommendation.

Expert consensus statements

Expert consensus statement:

12/12 experts concluded that in patients with asymptomatic carotid stenosis in whom revascularisation is considered to be appropriate and who are less suitable for surgery, stenting may be suggested. We recommend careful consideration of the risks and benefits at a multi-disciplinary team meeting.

Expert consensus statement:


12/12 experts concluded that the independently assessed risk of in-hospital stroke or death following endarterectomy or stenting for asymptomatic carotid stenosis should be as low as possible, ideally below 2%.(6)

Table 4. Summary of findings for stenting versus endarterectomy for asymptomatic carotid stenosis (PICO 3.1-3.11)


Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Updated Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		
PICO 3.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death												
6	randomised trials	not serious	not serious	not serious	serious ^a	none	86/2018 (4.3%)	46/1292 (3.6%)	RR 1.25 (0.88 to 1.79)	9 more per 1,000 (from 4 fewer to 28 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 3.2: Long-term risk of post-procedural ipsilateral stroke												
5	randomised trials	not serious	not serious	not serious	very serious ^b	none	23/926 (2.5%)	20/927 (2.2%)	RR 1.12 (0.62 to 2.00)	3 more per 1,000 (from 8 fewer to 22 more)	⊕⊕○○ LOW	CRITICAL
PICO 3.3: Long-term risk of stroke in any territory, including peri-procedural death												
5	randomised trials	not serious	not serious	not serious	serious ^a	none	68/929 (7.3%)	55/928 (5.9%)	RR 1.22 (0.87 to 1.71)	13 more per 1,000 (from 8 fewer to 42 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 3.4: Long-term risk of major stroke, including peri-procedural death												
3	randomised trials	not serious	not serious	not serious	very serious ^b	none	2/267 (0.7%)	2/273 (0.7%)	RR 0.99 (0.15 to 6.68)	0 fewer per 1,000 (from 20 fewer to 20 more)	⊕⊕○○ LOW	CRITICAL
PICO 3.5: Long-term risk of death												
4	randomised trials	not serious	not serious	not serious	very serious ^b	none	7/332 (2.1%)	9/340 (2.6%)	RR 0.82 (0.31 to 2.20)	5 fewer per 1,000 (from 18 fewer to 32 more)	⊕⊕○○ LOW	IMPORTANT
PICO 3.6: Risk of peri-procedural stroke												
7	randomised trials	not serious	not serious	not serious	serious ^a	none	52/2056 (2.5%)	19/1317 (1.4%)	RR 1.70 (0.99 to 2.93)	10 more per 1,000 (from 0 fewer to 28 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 3.7: Risk of peri-procedural death												
6	randomised trials	not serious	not serious	not serious	not serious	none	1/1462 (0.1%)	1/730 (0.1%)	RR 0.33 (0.02 to 5.33)	1 fewer per 1,000 (from 1 fewer to 6 more)	⊕⊕⊕⊕ HIGH	CRITICAL

Certainty assessment							No of patients		Effect		Certainty	Importance
No of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Updated Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		


PICO 3.8: Risk of peri-procedural stroke or death

7	randomised trials	not serious	not serious	not serious	serious ^a	none	53/2058 (2.6%)	20/1320 (1.5%)	RR 1.62 (0.96 to 2.76)	9 more per 1,000 (from 1 fewer to 27 more)	 MODERATE	CRITICAL
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PICO 3.9: Risk of peri-procedural major stroke or death

5	randomised trials	not serious	not serious	not serious	serious ^a	none	8/1776 (0.5%)	3/1033 (0.3%)	RR 1.54 (0.39 to 6.07)	2 more per 1,000 (from 2 fewer to 15 more)	 MODERATE	CRITICAL
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PICO 3.10: Risk of peri-procedural myocardial infarction

7	randomised trials	not serious	not serious	serious ^d	serious ^a	none	12/2041 (0.6%)	16/1304 (1.2%)	RR 0.53 (0.25 to 1.15)	6 fewer per 1,000 (from 9 fewer to 2 more)	 LOW	IMPORTANT
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PICO 3.11: Risk of peri-procedural cranial nerve injury

5	randomised trials	not serious	not serious	not serious	not serious	very strong association ^a	2/1823 (0.1%)	36/1092 (3.3%)	RR 0.09 (0.03 to 0.28)	30 fewer per 1,000 (from 32 fewer to 24 fewer)	 HIGH	IMPORTANT
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CI: Confidence interval; RR: Risk ratio

Explanations

- a. Few events, Wide confidence intervals
- b. Very wide confidence intervals
- c. Significant heterogeneity, I² > 60%
- d. Definition of myocardial infarction differed across trials
- e. Very large effect

4. Endarterectomy or medical therapy for symptomatic carotid stenosis

Description of studies

There are three RCTs which randomly assigned patients with symptomatic carotid artery stenosis to CEA or medical therapy alone in a 1:1 ratio. The *North American Symptomatic Carotid Endarterectomy Trial (NASCET)* separately reported results in patients with severe (70-99%), moderate (50-69%) or mild (<50%) symptomatic carotid stenosis.(9) The first report in 1991 included outcomes in 659 patients with severe stenosis who had experienced a hemispheric or retinal transient ischaemic attack (TIA) or a non-disabling stroke within the 120 days before enrolment.(61) The second report in 1998 included outcomes in 858 patients with moderate stenosis and 1368 patients with mild stenosis with a transient ischaemic attack or non-disabling stroke within 180 days before study entry.(62) The 1998 report also provided long-term follow-up data for up to eight years in patients with severe stenosis included in the first report.

The *MRC European Carotid Surgery Trial (ECST)* reported results in 778 patients with severe (70-99%) and 374 patients with very mild (0-29%) symptomatic carotid stenosis in 1991,(63, 64) the results in 1599 patients with mild to moderate (30-69%) symptomatic carotid stenosis in 1996, and the final results with follow-up for up to eight years in all 3024 patients with symptomatic carotid stenosis in 1998.(65) Eligible patients had a non-disabling ischaemic stroke, TIA or retinal infarction attributable to the carotid stenosis in the preceding 6 months. In the publication from which data for the current guideline were extracted, degrees of stenosis had been re-measured according to the method used in the NASCET trial.(12)

The *Veterans Affairs Cooperative Studies Program (VACSP)* symptomatic carotid stenosis trial included 189 patients with >50% symptomatic carotid stenosis and followed them up for a maximum of 33 months.(66) Eligible patients had an ischaemic stroke, TIA or transient monocular blindness in the preceding 120 days. Results were reported in 1991.

The effects of treatment are presented with best medical therapy alone as the reference group. A summary of findings is provided in table 5.

PICO 4.1: In patients with symptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death?

The reduction in the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death, with endarterectomy is strongly dependent on the degree of the symptomatic stenosis and the time interval between the index neurological event and randomisation. There is very low quality evidence for a benefit of CEA if data from all symptomatic patients, regardless of the severity of their stenosis, are grouped and analysed together (RR 0.83, 95%CI 0.61-1.14; equivalent to 26 fewer events with CEA per 1000 patients, from 59 fewer to 21 more; figure 4.1). The level of evidence was additionally downgraded for inconsistency due to statistical heterogeneity between trials. Stratifying results by degree of stenosis, there is moderate evidence of a meaningful benefit of CEA in patients with 70-99% stenosis (RR 0.37, 0.27-0.50; 169 fewer events per 1000 patients, from 196 fewer to 134 fewer; figure 4.1.4); low quality evidence of potential benefit in an overall population of patients with 50-69% stenosis (RR 0.82, 0.58-1.15; 29 fewer events per 1000 patients, from 67 fewer to 24 more; see also subgroup analysis below); and no evidence of benefit amongst patients with <50% stenosis (RR 1.09, 0.64 to 1.85) or near-occlusion (RR 1.03, 0.57 to 1.84; very low grade evidence each). The interaction between degree of stenosis and the effect of CEA was significant ($p < 0.0001$).

The benefit of CEA in patients with $\geq 50\%$ stenosis was most pronounced amongst patients randomised within two weeks of the index neurological event (RR 0.41, 0.30-0.58, 174 fewer events per 1000 patients, from 206 fewer to 124 fewer, high quality evidence; figure 4.1.3), but benefit was still present up to 12 weeks ($p = 0.001$ for interaction with time).

An individual patient data meta-analysis of all three trials showed that the degree of stenosis and time since the last event modified the effect of CEA in an additive manner. There was a significant 14.8% (95% CI 6.2-23.4%) absolute reduction in the 5-year risk of ipsilateral carotid territory ischaemic stroke or any stroke or death within 30 days of CEA in patients with moderate (50-69%) stenosis who were randomised within 14 days of their index ischaemic event (data not included in SoF table or figure). (67)

There is no evidence that the benefit of CEA differs between men and women or with age (figure 4.1.1 and 4.2.2). Although the reduction in the combined outcome was not statistically significant in women (likely due to the low number of women included in the trials; $n = 832$), the statistical test for interaction was not significant.

PICO 4.2: In patients with symptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of stroke in any territory, including peri-procedural death?

Amongst patients with all degrees of stenosis combined, there is moderate quality of evidence that CEA reduced the long-term risk of stroke in any territory, including peri-procedural death, compared with medical therapy alone (RR 0.85, 95% CI 0.77-0.94; 32 fewer events per 1000 patients, from 49 fewer to 13 fewer; figure 4.2). The evidence for a beneficial effect of CEA was of moderate quality in patients with 70-99% stenosis (RR 0.48, 95% CI 0.29-0.81; 171 fewer events per 1000 patients, from 233 fewer to 62 fewer; figure 4.2.1) and in patients with 50-59% stenosis (RR 0.77, 95% CI 0.63-0.94; 55 fewer events per 1000 patients, from 88 fewer to 14 fewer). Comparing the number of events prevented between PICO 4.1 and PICO 4.2 within each stenosis category, it can be inferred that CEA mainly prevents ipsilateral stroke.

PICO 4.3: In patients with symptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of major stroke, including peri-procedural death?

Amongst patients with all degrees of stenosis combined, endarterectomy did not significantly reduce the long-term risk of major stroke, including peri-procedural death (RR 0.79, 95% CI 0.51-1.22; 12 fewer events per 1000 patients, from 27 fewer to 12 more; low quality evidence; figure 4.3). However, once again, the benefit of CEA varies according to the degree of stenosis. In patients with 70-99% stenosis, there is high quality evidence of benefit (RR 0.35, 95% CI 0.22-0.57; 79 fewer events per 1000 patients, from 95 fewer to 52 fewer; figure 4.3.1). Conversely, there was low quality evidence of potential benefit in patients with 50-69% stenosis (RR 0.73, 95% CI 0.41 to 1.27; 15 fewer events per 1000 patients, from 33 fewer to 15 more), low quality evidence of harm in patients with <50% stenosis (RR 1.24, 95% CI 0.82 to 1.87), and very low quality evidence of harm in patients with near occlusion (RR 1.33, 95% CI 0.35 to 5.08).

PICO 4.4: In patients with symptomatic carotid stenosis, does endarterectomy compared with medical therapy alone reduce the long-term risk of death?

Endarterectomy does not reduce the long-term risk of death compared with medical therapy alone (RR 1.03, 95% CI 0.65-1.65; 6 more events per 1000 patients, from 74 fewer to 137 more; very low quality of evidence; figure 4.4).

Analysis of current evidence and evidence-based recommendation

Evidence of the effect of CEA compared with medical therapy alone for symptomatic carotid stenosis was available from 3 trials, which included 6,098 patients. Symptomatic carotid stenosis was defined by the occurrence of ischaemic ocular or cerebral events attributable to the stenosis within four to six months before enrolment, depending on the trial and the severity of stenosis. The evidence provided relates to the time when these trials were performed three decades ago. Medical treatment of patients with atherosclerotic carotid stenosis has improved, with widespread use of statins, the availability of better antiplatelet treatment regimens and stricter control of blood pressure. However, surgical techniques and perioperative management have also improved since these trials were completed. We therefore downgraded the overall quality of evidence for indirectness.

The benefits of CEA in patients with symptomatic carotid stenosis strongly depends on the degree of stenosis. Amongst patients with severe (70-99%) stenosis, there is moderate quality evidence that CEA prevents ipsilateral stroke and stroke in any territory, and high-grade evidence that it prevents major stroke, taking into account the combined risks of peri-operative stroke or death. In patients with moderate (50-69%) carotid stenosis, there is low quality evidence that CEA prevents ipsilateral stroke and major stroke, and moderate quality evidence for prevention of stroke in any territory, again taking into account the peri-operative stroke or death risk, if patients are operated upon within 14 days of their presenting cerebrovascular event. There is no evidence that CEA prevents stroke in patients with mild (<50%) stenosis or near-occlusion of the carotid artery. However, the definition of near-occlusion in the early endarterectomy trials depended on intra-arterial angiography, and there are no widely-accepted standardised criteria for near-occlusion on Doppler ultrasound or non-invasive angiography (68) We therefore could not make any clear recommendations on the treatment of carotid near-occlusion in this guideline. The benefit of CEA also strongly depends on the timing of treatment, with the greatest reduction in stroke risk achieved if surgery is performed <14 days of the index event. We found no evidence that the benefit of CEA varies significantly between men and women or between older and younger patients.

The optimal management of patients with distal tandem stenosis is uncertain. In NASCET, patients who had 85-99% extracranial ICA stenosis and any degree of co-existing, ipsilateral intracranial atherosclerotic disease (IAD) had an increased risk of ipsilateral stroke over 3 years if they were treated with best medical therapy alone compared with those without IAD (45.7% vs. 25.3%, relative risk 1.8; 95% CI: 1.1 – 3.2).(69) However, the three-year risk of ipsilateral stroke in surgically-treated patients with 85-99% extracranial ICA stenosis was similar in those with and those without IAD (8.6%

vs. 10%, relative risk 0.9; 95% CI: 0.2 – 3.0). Therefore, IAD should not deter one from proceeding to CEA in suitable patients, whilst acknowledging that only a very small number of patients with severe stenosis were included in this subgroup analysis of the NASCET data.

Recommendations:

In patients with severe (70-99%) symptomatic carotid artery stenosis, we recommend carotid endarterectomy.

Quality of evidence: **Moderate** ⊕⊕⊕

Strength of recommendation: **Strong for carotid endarterectomy** ↑↑

In patients with moderate (50-69%) symptomatic carotid artery stenosis, we suggest carotid endarterectomy.

Quality of evidence: **Low** ⊕⊕

Strength of recommendation: **Weak for carotid endarterectomy** ↑

In patients with mild (<50%) symptomatic carotid artery stenosis, we recommend against carotid endarterectomy.

Quality of evidence: **Very low** ⊕

Strength of recommendation: **Strong against carotid endarterectomy** ↓↓

In patients with 50-99% symptomatic carotid stenosis in whom surgery is considered appropriate, we recommend early endarterectomy, ideally within two weeks of the first neurological event.

Quality of evidence: **High** ⊕⊕⊕⊕







Strength of recommendation: **Strong for carotid endarterectomy** ↑↑

These recommendations are independent of sex and age.

Additional information

The *Second European Carotid Surgery Trial (ECST-2)* is comparing OMT alone versus OMT and carotid revascularisation in patients with symptomatic carotid stenosis estimated to be at low or intermediate risk of stroke using 'clinical risk modelling', and in patients with asymptomatic carotid stenosis. The ECST-2 trial was prematurely discontinued after inclusion of 429 patients due to slow recruitment and results are awaited (www.ecst2.com, last access Feb 2, 2021).

Table 5. Summary of findings for endarterectomy versus medical therapy for symptomatic carotid stenosis (PICO 4.1-4.4).

Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 4.1: Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		
PICO 4.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death												
3	randomised trials	not serious	serious ^a	serious ^b	serious ^c	none	394/3336 (11.8%)	415/2754 (15.1%)	RR 0.83 (0.61 to 1.14)	26 fewer per 1,000 (from 59 fewer to 21 more)	 VERY LOW	CRITICAL
PICO 4.1.1a: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Age < 65 years												
2	randomised trials	not serious	not serious	serious ^b	not serious	none	86/731 (11.8%)	93/550 (16.9%)	RR 0.70 (0.53 to 0.92)	51 fewer per 1,000 (from 79 fewer to 14 fewer)	 MODERATE	CRITICAL
PICO 4.1.1b: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Age ≥ 65 years												
2	randomised trials	not serious	not serious	serious ^b	not serious	none	89/743 (12.0%)	151/694 (21.8%)	RR 0.57 (0.44 to 0.73)	94 fewer per 1,000 (from 122 fewer to 59 fewer)	 MODERATE	CRITICAL
PICO 4.1.2a: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Men												
2	randomised trials	not serious	not serious	serious ^b	not serious	none	112/1013 (11.1%)	184/873 (21.1%)	RR 0.54 (0.44 to 0.67)	97 fewer per 1,000 (from 118 fewer to 70 fewer)	 MODERATE	CRITICAL
PICO 4.1.2.b.: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Women												
2	randomised trials	not serious	not serious	serious ^b	serious ^c	none	63/461 (13.7%)	60/371 (16.2%)	RR 0.85 (0.58 to 1.23)	24 fewer per 1,000 (from 68 fewer to 37 more)	 LOW	CRITICAL
PICO 4.1.3a: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: <2 weeks since most recent ischaemic event												
2	randomised trials	not serious	not serious	serious ^b	not serious	strong association	40/325 (12.3%)	88/299 (29.4%)	RR 0.41 (0.30 to 0.58)	174 fewer per 1,000 (from 206 fewer to 124 fewer)	 HIGH	CRITICAL

Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 4.1: Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		
PICO 4.1.3b: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: 2-4 weeks since most recent ischaemic event												
2	randomised trials	not serious	not serious	serious ^b	not serious	none	31/268 (11.6%)	44/215 (20.5%)	RR 0.58 (0.35 to 0.98)	86 fewer per 1,000 (from 133 fewer to 4 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 4.1.3c: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: 4-12 weeks since most recent ischaemic event												
2	randomised trials	not serious	not serious	serious ^b	not serious	none	63/560 (11.3%)	81/498 (16.3%)	RR 0.70 (0.51 to 0.95)	49 fewer per 1,000 (from 80 fewer to 8 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 4.1.3d: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: >12 weeks since most recent ischaemic event												
2	randomised trials	not serious	not serious	serious ^b	very serious ^d	none	41/321 (12.8%)	31/232 (13.4%)	RR 1.01 (0.66 to 1.57)	1 more per 1,000 (from 45 fewer to 76 more)	⊕○○○ VERY LOW	CRITICAL
PICO 4.1.4a: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Near occlusion												
2	randomised trials	not serious	not serious	serious ^b	very serious ^d	none	24/157 (15.3%)	17/114 (14.9%)	RR 1.03 (0.57 to 1.84)	4 more per 1,000 (from 64 fewer to 125 more)	⊕○○○ VERY LOW	CRITICAL
PICO 4.1.4b: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Severe (70-99%) stenosis												
2	randomised trials	not serious	serious ^a	serious ^b	not serious	strong association	50/518 (9.7%)	117/436 (26.8%)	RR 0.37 (0.27 to 0.50)	169 fewer per 1,000 (from 196 fewer to 134 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 4.1.4c: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Moderate (50-69%) stenosis												
2	randomised trials	not serious	not serious	serious ^b	serious ^c	none	101/808 (12.5%)	110/694 (15.9%)	RR 0.82 (0.58 to 1.15)	29 fewer per 1,000 (from 67 fewer to 24 more)	⊕⊕○○ LOW	CRITICAL
PICO 4.1.4d: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death: Mild (<50%) stenosis												
2	randomised trials	not serious	serious ^a	serious ^b	serious ^c	none	212/1762 (12.0%)	164/1413 (11.6%)	RR 1.09 (0.64 to 1.85)	10 more per 1,000 (from 42 fewer to 99 more)	⊕○○○ VERY LOW	CRITICAL

Certainty assessment							N ^e of patients		Effect		Certainty	Importance
N ^e of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 4.1: Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		

PICO 4.2: Long-term risk of stroke in any territory, including peri-procedural death

3	randomised trials	not serious	not serious	serious ^b	not serious	none	586/3336 (17.6%)	584/2754 (21.2%)	RR 0.85 (0.77 to 0.94)	32 fewer per 1,000 (from 49 fewer to 13 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
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PICO 4.2.1a: Long-term risk of stroke in any territory, including peri-procedural death: Near-occlusion

2	randomised trials	not serious	not serious	serious ^b	very serious ^d	none	32/157 (20.4%)	25/114 (21.9%)	RR 1.00 (0.46 to 2.21)	0 fewer per 1,000 (from 118 fewer to 265 more)	⊕○○○ VERY LOW	CRITICAL
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PICO 4.2.1b: Long-term risk of stroke in any territory, including peri-procedural death: Severe (70-99%) stenosis

2	randomised trials	not serious	serious ^a	serious ^b	not serious	strong association ^d	84/518 (16.2%)	143/436 (32.8%)	RR 0.48 (0.29 to 0.81)	171 fewer per 1,000 (from 233 fewer to 62 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
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PICO 4.2.1c: Long-term risk of stroke in any territory, including peri-procedural death: Moderate (50-69%) stenosis

2	randomised trials	not serious	not serious	serious ^b	not serious	none	144/808 (17.8%)	165/694 (23.8%)	RR 0.77 (0.63 to 0.94)	55 fewer per 1,000 (from 88 fewer to 14 fewer)	⊕⊕⊕○ MODERATE	CRITICAL
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PICO 4.2.1d: Long-term risk of stroke in any territory, including peri-procedural death: Mild (<50%) stenosis

2	randomised trials	not serious	not serious	serious ^b	serious ^c	none	318/1762 (18.0%)	244/1413 (17.3%)	RR 1.09 (0.89 to 1.34)	16 more per 1,000 (from 19 fewer to 59 more)	⊕⊕○○ LOW	CRITICAL
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PICO 4.3: Long-term risk of major stroke, including peri-procedural death

3	randomised trials	not serious	not serious	serious ^b	serious ^c	none	150/3336 (4.5%)	152/2754 (5.5%)	RR 0.79 (0.51 to 1.22)	12 fewer per 1,000 (from 27 fewer to 12 more)	⊕⊕○○ LOW	CRITICAL
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PICO 4.3.1a: Long-term risk of major stroke, including peri-procedural death: Near-occlusion

2	randomised trials	not serious	not serious	serious ^b	very serious ^d	none	12/157 (7.6%)	7/114 (6.1%)	RR 1.33 (0.35 to 5.08)	20 more per 1,000 (from 40 fewer to 251 more)	⊕○○○ VERY LOW	CRITICAL
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Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 4.1: Endarterectomy	Medical therapy	Relative (95% CI)	Absolute (95% CI)		
PICO 4.3.1b: Long-term risk of major stroke, including peri-procedural death: Severe (70-99%) stenosis												
2	randomised trials	not serious	not serious	serious ^b	not serious	strong association ^f	22/518 (4.2%)	53/436 (12.2%)	RR 0.35 (0.22 to 0.57)	79 fewer per 1,000 (from 95 fewer to 52 fewer)	⊕⊕⊕⊕ HIGH	CRITICAL
PICO 4.3.1c: Long-term risk of major stroke, including peri-procedural death: Moderate (50-69%) stenosis												
2	randomised trials	not serious	not serious	serious ^b	serious ^c	none	35/808 (4.3%)	39/694 (5.6%)	RR 0.73 (0.41 to 1.27)	15 fewer per 1,000 (from 33 fewer to 15 more)	⊕⊕○○ LOW	CRITICAL
PICO 4.3.1d: Long-term risk of major stroke, including peri-procedural death: Mild (<50%) stenosis												
2	randomised trials	not serious	not serious	serious ^b	serious ^c	none	79/1762 (4.5%)	50/1413 (3.5%)	RR 1.24 (0.82 to 1.87)	8 more per 1,000 (from 6 fewer to 31 more)	⊕⊕○○ LOW	CRITICAL
PICO 4.4: Long term risk of death												
3	randomised trials	not serious	not serious	serious ^b	very serious ^d	none	520/2227 (23.3%)	345/1640 (21.0%)	RR 1.03 (0.65 to 1.65)	6 more per 1,000 (from 74 fewer to 137 more)	⊕○○○ VERY LOW	IMPORTANT

CI: Confidence interval; RR: Risk ratio

Explanations

- Significant heterogeneity, $I^2 > 70\%$
- Endarterectomy and medical therapy have evolved since the trials contributing the evidence were performed
- Few events, wide confidence intervals
- Very wide confidence intervals
- Significant heterogeneity, $I^2 = 68\%$, 69%
- Large effect

5. Stenting or medical therapy for symptomatic carotid stenosis

Description of studies

We identified no RCTs comparing stenting versus medical therapy alone in patients with symptomatic carotid stenosis that fulfilled our inclusion criteria. We excluded two small RCTs because these studies did not report outcomes according to symptom status,(38, 39) or patients were treated with primary balloon angioplasty.(39)

6. Stenting or endarterectomy for symptomatic carotid stenosis

Description of studies

A single-centre trial in Lexington, Kentucky, USA randomised 104 patients with $\geq 70\%$ symptomatic carotid stenosis to receive either CAS without a cerebral protection device (CPD) or CEA and reported results up to 2 years after randomisation in 2001.(70)

The French multi-centre *Endarterectomy versus Angioplasty in Patients with Symptomatic Severe Carotid Stenosis (EVA-3S)* trial randomised 527 patients with $\geq 60\%$ symptomatic carotid stenosis to undergo CAS or CEA between 2000-2005.(71-76) Interventionists were required to have performed at least 12 CAS procedures, or at least 35 stenting procedures in the supra-aortic trunks, of which at least 5 involved the carotid artery. The use of CPDs during stenting was made mandatory after an interim analysis raised safety concerns amongst patients treated without CPDs. The trial was stopped early for safety and futility reasons. Initial results were published in 2006, and final results with available data over a median follow-up period of 7.1 years were reported in 2014.

The multi-centre *Stent-supported Percutaneous Angioplasty of the Carotid artery versus Endarterectomy (SPACE)* trial randomised 1214 patients with $\geq 50\%$ symptomatic carotid stenosis between CAS and CEA in Germany, Austria, and Switzerland between 2001-2006.(77-79) Interventionists had to show proof of at least 25 successful, consecutive percutaneous transluminal angioplasty or stent procedures in the carotid artery. The use of a CPD was not mandatory. The trial was stopped early for reasons of futility and lack of funding. Initial results were published in 2006 and final results up to two years after randomization were published in 2008.

A single-centre trial in Regensburg, Germany, randomized 87 patients with $\geq 70\%$ symptomatic carotid stenosis to undergo CAS without a CPD or CEA between 1999- 2002.(80) Recruitment was stopped when the multi-centre *SPACE* trial, which had a similar study design, was commenced. Results over a median follow-up period of >5 years were published in 2008.

The multi-centre *International Carotid Stenting Study (ICSS)* randomised 1713 patients with $\geq 50\%$ symptomatic carotid stenosis to receive either CAS or CEA in Europe, Australia, New Zealand, and Canada between 2001 and 2008.(81-84) Eligible patients had symptoms attributable to their carotid stenosis within 12 months before randomisation; however, only 4% had symptoms which occurred more than 6 months before randomisation. Interventionists were required to have carried out at least 50 stenting procedures, at least 10 of which were in the carotid artery. Use of CPDs was recommended

but not mandatory. Initial results were published in 2011 and final results with data over a median follow-up period of 4.2 years were reported in 2015.

The single-centre *Basel Carotid Artery Stenting Study (BACASS)* randomised 20 patients with $\geq 50\%$ symptomatic carotid stenosis to CAS with routine use of a CPD or CEA between 1998-2002.(85) Recruitment was stopped when the centre started recruiting patients in ICSS. Results including follow-up data over a median of four years after randomisation were published in 2008.

We also extracted relevant outcomes in symptomatic patients from the *Czech Republic* and *CREST* trials, which are described in results section 3. Furthermore, we included outcomes in pre-defined patient subgroups derived from pooled analyses of individual patient data (IPD) from the *EVA-3S*, *SPACE*, *ICSS* and *CREST* trials which were performed by the *Carotid Stenosis Trialists' Collaboration (CSTC)*.(86-88)

We excluded one industry-funded multi-centre randomised trial because the results were only reported in a conference abstract,(89) and also excluded one single-centre and one multicentre randomised trial in which the majority of patients in the endovascular group were treated with primary balloon angioplasty.(90, 91)

The effects of treatment are presented with endarterectomy as the reference group. A summary of findings is provided in table 6.

PICO 6.1: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death?

There is moderate quality of evidence that endarterectomy is superior to stenting in preventing the combined outcome of post-procedural ipsilateral stroke, peri-procedural stroke in any territory, or peri-procedural death (RR 1.43, 95% CI 1.17-1.75; equivalent to 31 more events with stenting per 1000 patients, from 12 more to 54 more; figure 6.1). In a pooled IPD analysis from the *EVA-3S*, *SPACE*, *ICSS* and *CREST* trials, the relative risk of this outcome varied with age:(88) this analysis provided moderate quality evidence that CEA was superior to CAS in patients aged 65-74 years (hazard ratio [HR] 1.67, 95% CI 1.23-2.27) and ≥ 75 years (HR 1.85, 95% CI 1.35-2.53), and low quality evidence that there was no difference in outcomes between stenting and endarterectomy amongst patients < 65 years old (HR 0.83, 95% CI 0.56-1.21), with a significant interaction between age and treatment effect ($p=0.003$; data not shown in figure). There was no evidence of an interaction with sex or severity of stenosis.

PICO 6.2: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of post-procedural ipsilateral stroke?

There is moderate quality of evidence that stenting and endarterectomy do not differ in their ability to prevent long-term post-procedural ipsilateral stroke (RR 1.06, 95% CI 0.74-1.51; equivalent to 1 more event with stenting per 1000 patients, from 6 fewer to 12 more; figure 6.2).

PICO 6.3: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of stroke in any territory, including peri-procedural death?

There is moderate quality of evidence that endarterectomy is superior to stenting in preventing the combined long-term outcome of stroke in any territory or peri-procedural death (RR 1.34, 95% CI 1.08-1.66; 35 more events with stenting per 1000 patients, from 8 more to 68 more; figure 6.3).

PICO 6.4: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of major stroke, including peri-procedural death?

There is low quality of evidence that endarterectomy and stenting do not differ in the long-term risk of major stroke or peri-procedural death (RR 1.19, 95% CI 0.88-1.62; 12 more events with stenting per 1000 patients, from 8 fewer to 39 more; figure 6.4).

PICO 6.5: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of death?

There is low quality of evidence that endarterectomy and stenting do not differ in the long-term risk of death (RR 1.09, 95% CI 0.94-1.27; 13 more events with stenting per 1000 patients, from 9 fewer to 38 more; figure 6.5).

PICO 6.6: In patients with asymptomatic or symptomatic carotid stenosis, do endarterectomy and stenting differ in the long-term risk of severe restenosis?

For the analysis of restenosis, we combined the data from trials including patients with asymptomatic carotid stenosis, symptomatic stenosis, or both. There is very low quality evidence that endarterectomy and stenting do not differ in the long-term risk of severe restenosis (RR 1.34, 95% CI 0.88-2.03; figure 6.6). We additionally downgraded the evidence for inconsistency, as there was evidence of substantial heterogeneity between trials ($I^2=57\%$).

PICO 6.7: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural stroke?

There is moderate quality of evidence that stenting is associated with a higher risk of peri-procedural stroke than endarterectomy (RR 1.64, 95% CI 1.24-2.17; 26 more events with stenting per 1000 patients, from 10 more to 48 more; figure 6.7).

PICO 6.8: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural death?

There is very low quality of evidence that stenting and endarterectomy do not differ in the risk of peri-procedural death (RR 1.45, 95% CI 0.73-2.87; 3 more events per 1000 patients with stenting, from 2 fewer to 10 more; figure 6.8).

PICO 6.9: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural stroke or death?

There is moderate quality evidence that stenting is associated with a higher risk of peri-procedural stroke or death than endarterectomy overall (RR 1.68, 95% CI 1.20-2.34; 28 more events with stenting per 1000 patients, from 8 more to 55 more; figure 6.9). However, these results vary with age. Amongst patients ≥ 70 years, there is high quality evidence that CAS is associated with a higher risk of this composite outcome compared with CEA (RR 2.10, 95% CI 1.55-2.84; 53 more events with stenting per 1000 patients, from 26 more to 88 more; figure 6.9.1). Amongst patients < 70 years, there is low quality evidence that the risk of this combined outcome does not differ between the two treatment modalities (RR 1.10, 95% CI 0.75-1.60; 4 more events with stenting per 1000 patients, from 10 fewer to 24 more). The interaction between age and treatment effect is significant ($p=0.009$), but there is no evidence of an interaction with sex (figure 6.9.2).

A pooled analysis of IPD from *EVA-3S*, *SPACE* and *ICSS* provides no evidence for a modification of the effect of CAS versus CEA on the risk of peri-procedural stroke or death by the severity of stenosis (figure 6.9.3) or type of most recent ischaemic event (hemispheric stroke, transient ischaemic attack or ocular ischaemia; figure 6.9.5).⁽⁸⁶⁾

Another pooled analysis of IPD from *EVA-3S*, *SPACE*, *ICSS* and *CREST* provides high-quality evidence of an increased risk of peri-procedural stroke or death with CAS compared with CEA amongst patients treated < 7 days after their most recent ischaemic event (RR 6.30, 95% CI 1.92-20.66; 70 more events with CAS per 1000 patients, from 12 more to 261 more; figure 6.9.4), and moderate quality evidence

for this difference amongst patients treated >7 days after the event (RR 2.00, 95% CI 1.50-2.68; 36 more events with stenting per 1000 patients, from 18 more to 60 more).(87) The unadjusted p-value for the interaction between timing and treatment effect was 0.07, the adjusted p-value in the original publication was 0.06.

PICO 6.10: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural major stroke or death?

There is low quality of evidence that endarterectomy and stenting do not differ in the risk of peri-procedural major stroke or death (RR 1.33, 95% CI 0.96-1.85; 8 more events with stenting per 1000 patients, from 1 fewer to 21 more; figure 6.10).

PICO 6.11: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural myocardial infarction?

There is moderate quality of evidence that stenting is associated with a lower risk of peri-procedural myocardial infarction than endarterectomy (RR 0.48, 95% CI 0.24-0.98; 6 fewer events per 1000, from 9 fewer to 0 fewer; figure 6.11). Even though the relative effect was large, there were a limited number of clinically relevant cardiac outcome events observed. Furthermore, we had additional concerns about ‘indirectness’ due to the definition of myocardial infarction used in the CREST trial which contributed to two thirds of the cardiac outcome events included in the aggregate analysis (see results section 3).

PICO 6.12: In patients with symptomatic carotid stenosis, do endarterectomy and stenting differ in the risk of peri-procedural cranial nerve injury?

There is strong evidence that stenting is associated with a lower risk of peri-procedural cranial nerve injury than endarterectomy (RR 0.10, 95% CI 0.05-0.20; 49 fewer events with stenting per 1000 patients, from 52 fewer to 44 fewer; figure 6.12). We upgraded the quality of the evidence by 2 levels for strength of effect.

Analysis of current evidence and evidence-based recommendation

Evidence to compare short-term risks and long-term effects of CAS versus CEA for the treatment of symptomatic carotid stenosis was derived from 7 RCTs which included a total of 4893 patients. It is important to note that the available evidence for CAS relates to percutaneous trans-femoral stenting only. There are no available data from RCTs on the safety of trans-carotid stenting. As such, all recommendations included in this guideline refer to trans-femoral CAS. All studies included patients

with $\geq 50\%$ stenosis. Symptomatic carotid stenosis was defined by the occurrence of ischaemic ocular or cerebral events attributable to the stenosis within six months prior to enrolment, except in ICSS where a very small minority of patients were enrolled 6-12 months after symptom onset. Amongst the four largest trials contributing to the evidence, the median duration of follow-up was four to seven years in three studies and two years in one study. When recruitment in these trials started 20 years ago, carotid artery stenting was still at a relatively early stage of technical development, peri-procedural medication regimens were not standardised, and there was limited experience with the procedure. In addition, only a minority of patients included in these trials were treated within the recommended 14 days of their index ischaemic event. We therefore downgraded the quality of evidence for indirectness.

Overall, there is moderate quality evidence that endarterectomy is superior to stenting when one considers peri-procedural and post-procedural outcomes that were rated as ‘critical’ for decision making. The differences between stenting and endarterectomy are mainly apparent in the peri-procedural period: stenting is associated with a higher risk of peri-procedural stroke than endarterectomy (critical for decision making), whereas endarterectomy is associated with higher risks of myocardial infarction and mostly transient cranial nerve palsy (important for decision making).

The risks of peri-procedural stroke or death differ between patient subgroups: there is high quality evidence that stenting is associated with a higher risk of this outcome in patients ≥ 70 years, and low quality evidence that the risk of this outcome is similar in patients < 70 years. The higher risk of peri-procedural stroke or death after carotid artery stenting compared with endarterectomy is also more evident amongst patients treated within 7 days of their index cerebrovascular event. After the peri-procedural period, there is moderate grade evidence that stenting and endarterectomy do not differ in their ability to prevent stroke.

Recommendations

In patients with symptomatic carotid artery stenosis requiring revascularisation, we recommend endarterectomy as the treatment of choice.

Quality of evidence: Moderate ⊕⊕⊕

Strength of recommendation: Strong for carotid endarterectomy ↑↑

In patients with symptomatic carotid stenosis <70 years old requiring revascularisation, we suggest that stenting may be considered as an alternative to endarterectomy.

Quality of evidence: Low ⊕⊕

Strength of recommendation: Weak for carotid stenting ↑

Additional information

In light of technical developments in stent design and cerebral protection devices, and alternative (trans-brachial and trans-carotid) access routes which are now available, new trials of stenting in selected patients with symptomatic carotid stenosis are warranted.

Expert consensus statements

12/12 experts concluded that the suitability of a patient with symptomatic carotid stenosis for carotid endarterectomy versus stenting should also take into account the interval since their last ischaemic cerebrovascular event, as well as anatomical and morphological features, including the atherosclerotic burden of the aortic arch.

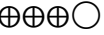
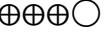


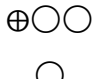
11/12 experts concluded that the independently assessed risk of in-hospital stroke or death following endarterectomy or stenting for symptomatic carotid stenosis should not exceed 4%.(6)

12/12 experts concluded that where possible, the indication for carotid endarterectomy or carotid artery stenting should be discussed at a multi-disciplinary team meeting. Consensus decisions can be made in between meetings, in order not to delay urgent revascularisations.

12/12 experts concluded that the establishment of validated local, regional or national registries, including audit systems for carotid interventions to monitor complication rates in patients with asymptomatic and symptomatic carotid stenosis is recommended.

Table 6. Summary of findings for stenting or endarterectomy for symptomatic carotid stenosis (PICO 6.1-6.12).


Certainty assessment							Nº of patients		Effect		Certainty	Importance
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		
PICO 6.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death												
7	randomised trials	not serious	not serious	serious ^a	not serious	none	261/2499 (10.4%)	177/2466 (7.2%)	RR 1.43 (1.17 to 1.75)	31 more per 1,000 (from 12 more to 54 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.1.1a: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Age < 65 years												
4	IPD of randomised trials	not serious	not serious	serious ^a	serious ^b	none			HR 0.83 (0.56 to 1.21)		⊕⊕○○ LOW	CRITICAL
PICO 6.1.1b: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Age 65-74 years												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none			HR 1.67 (1.23 to 2.27)		⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.1.1c: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Age ≥ 75 years												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none			HR 1.85 (1.35 to 2.53)		⊕⊕⊕○ MODERATE	CRITICAL
6.1.2a: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Men												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none			HR 1.54 (1.23 to 1.95)		⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.1.2b: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Women												
4	IPD of randomised trials	not serious	not serious	serious ^a	serious ^b	none			HR 1.25 (0.90 to 1.74)		⊕⊕○○ LOW	CRITICAL
PICO 6.1.3a: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Severe stenosis												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none			HR 1.48 (1.20 to 1.81)		⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.1.3b: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death: Moderate stenosis												
4	IPD of randomised trials	not serious	not serious	serious ^a	serious ^b	none			HR 1.33 (0.84 to 2.10)		⊕⊕○○ LOW	CRITICAL

Certainty assessment							N° of patients		Effect		Certainty	Importance
N° of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		
PICO 6.2: Long-term risk of post-procedural ipsilateral stroke												
6	randomised trials	not serious	not serious	serious ^a	not serious	none	62/2429 (2.6%)	58/2408 (2.4%)	RR 1.06 (0.74 to 1.51)	1 more per 1,000 (from 6 fewer to 12 more)	 MODERATE	CRITICAL
PICO 6.3. Long-term risk of stroke in any territory, including peri-procedural death												
6	randomised trials	not serious	not serious	serious ^a	not serious	none	352/2435 (14.5%)	247/2411 (10.2%)	RR 1.34 (1.08 to 1.66)	35 more per 1,000 (from 8 more to 68 more)	 MODERATE	CRITICAL
PICO 6.4. Long-term risk of major stroke and peri-procedural death												
3	randomised trials	not serious	not serious	serious ^a	serious ^b	none	84/1117 (7.5%)	71/1125 (6.3%)	RR 1.19 (0.88 to 1.62)	12 more per 1,000 (from 8 fewer to 39 more)	 LOW	CRITICAL
PICO 6.5. Long-term risk of death												
5	randomised trials	not serious	not serious	serious ^a	serious ^b	none	278/1778 (15.6%)	251/1762 (14.2%)	RR 1.09 (0.94 to 1.27)	13 more per 1,000 (from 9 fewer to 38 more)	 LOW	IMPORTANT
PICO 6.6: Long-term risk of severe restenosis												
9	randomised trials	not serious	serious ^c	serious ^a	serious ^b	none	212/3077 (6.9%)	166/3147 (5.3%)	RR 1.37 (0.89 to 2.10)	20 more per 1,000 (from 6 fewer to 58 more)	 VERY LOW	IMPORTANT
PICO 6.7: Risk of peri-procedural stroke												


Certainty assessment							Nº of patients		Effect		Certainty	Importance
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		
7	randomised trials	not serious	not serious	serious ^a	not serious	none	168/249 5 (6.7%)	101/2472 (4.1%)	RR 1.64 (1.24 to 2.17)	26 more per 1,000 (from 10 more to 48 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.8: Risk of peri-procedural death												
8	randomised trials	not serious	not serious	serious ^a	very serious ^d	none	22/2538 (0.9%)	14/2514 (0.6%)	RR 1.45 (0.73 to 2.87)	3 more per 1,000 (from 2 fewer to 10 more)	⊕○○○ ○ VERY LOW	CRITICAL
PICO 6.9: Risk of peri-procedural stroke or death												
7	randomised trials	not serious	not serious	serious ^a	not serious	none	172/249 5 (6.9%)	101/2470 (4.1%)	RR 1.68 (1.20 to 2.34)	28 more per 1,000 (from 8 more to 55 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.9.1a: Risk of peri-procedural stroke or death: Age < 70 years												
6	randomised trials	not serious	not serious	serious ^a	serious ^b	none	56/1247 (4.5%)	49/1206 (4.1%)	RR 1.10 (0.75 to 1.60)	4 more per 1,000 (from 10 fewer to 24 more)	⊕⊕○○ LOW	CRITICAL
PICO 6.9.1b: Risk of peri-procedural stroke or death: Age ≥ 70 years												
6	randomised trials	not serious	not serious	serious ^a	not serious	strong association ^e	122/1195 5 (10.2%)	58/1213 (4.8%)	RR 2.10 (1.55 to 2.84)	53 more per 1,000 (from 26 more to 88 more)	⊕⊕⊕⊕ HIGH	CRITICAL

Certainty assessment							No of patients		Effect		Certainty	Importance
No of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		


PICO 6.9.2a: Risk of peri-procedural stroke or death: Men

6	randomised trials	not serious	not serious	serious ^a	not serious	none	121/1695 (7.1%)	70/1700 (4.1%)	RR 1.76 (1.09 to 2.85)	31 more per 1,000 (from 4 more to 76 more)	 MODERATE	CRITICAL
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PICO 6.9.2b: Risk of peri-procedural stroke or death: Women

6	randomised trials	not serious	not serious	serious ^a	serious ^b	none	57/747 (7.6%)	36/719 (5.0%)	RR 1.45 (0.94 to 2.23)	23 more per 1,000 (from 3 fewer to 62 more)	 LOW	CRITICAL
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PICO 6.9.3a: Risk of peri-procedural stroke or death: Severe (70-99%) stenosis

3	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none	132/1393 (9.5%)	86/1381 (6.2%)	RR 1.52 (1.17 to 1.98)	32 more per 1,000 (from 11 more to 61 more)	 MODERATE	CRITICAL
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
PICO 6.9.3b: Risk of peri-procedural stroke or death: Moderate (50-69%) stenosis

3	IPD of randomised trials	not serious	not serious	serious ^a	serious ^d	none	21/332 (6.3%)	13/327 (4.0%)	RR 1.59 (0.81 to 3.12)	23 more per 1,000 (from 8 fewer to 84 more)	 LOW	CRITICAL
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
Certainty assessment							№ of patients		Effect		Certainty	Importance
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		
PICO 6.9.4a: Risk of peri-procedural stroke or death: ≤ 7 days since most recent ischaemic event												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	very strong association ^a	24/287 (8.4%)	3/226 (1.3%)	RR 6.30 (1.92 to 20.66)	70 more per 1,000 (from 12 more to 261 more)	⊕⊕⊕⊕ HIGH	CRITICAL
PICO 6.9.4b: Risk of peri-procedural stroke or death: > 7 days since most recent ischaemic event												
4	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none	129/1798 (7.2%)	65/1815 (3.6%)	RR 2.00 (1.50 to 2.68)	36 more per 1,000 (from 18 more to 60 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.9.5a: Risk of peri-procedural stroke or death: Hemispheric stroke as most recent ischaemic event												
3	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none	85/813 (10.5%)	52/797 (6.5%)	RR 1.60 (1.15 to 2.23)	39 more per 1,000 (from 10 more to 80 more)	⊕⊕⊕○ MODERATE	CRITICAL
PICO 6.9.5b: Risk of peri-procedural stroke or death: TIA as most recent ischaemic event												
3	IPD of randomised trials	not serious	not serious	serious ^a	not serious	none	53/589 (9.0%)	31/601 (5.2%)	RR 1.74 (1.14 to 2.68)	38 more per 1,000 (from 7 more to 87 more)	⊕⊕⊕○ MODERATE	CRITICAL

Certainty assessment							N _e of patients		Effect		Certainty	Importance
N _e of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	PICO 6.1: Stenting	Endarterectomy	Relative (95% CI)	Absolute (95% CI)		


PICO 6.9.5c: Risk of peri-procedural stroke or death: Retinal ischaemia as most recent ischaemic event

3	IPD of randomised trials	not serious	not serious	serious ^a	very serious ^d	none	15/310 (4.8%)	14/297 (4.7%)	RR 1.03 (0.50 to 2.09)	1 more per 1,000 (from 24 fewer to 51 more)	 VERY LOW	CRITICAL
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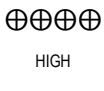
PICO 6.10: Risk of peri-procedural major stroke or death

7	randomised trials	not serious	not serious	serious ^a	serious ^b	none	81/2495 (3.2%)	60/2470 (2.4%)	RR 1.33 (0.96 to 1.85)	8 more per 1,000 (from 1 fewer to 21 more)	 LOW	
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PICO 6.11: Risk of peri-procedural myocardial infarction

6	randomised trials	not serious	not serious	serious ^a	serious ^b	strong association ^a	11/1878 (0.6%)	23/1874 (1.2%)	RR 0.48 (0.24 to 0.98)	6 fewer per 1,000 (from 9 fewer to 0 fewer)	 MODERATE	IMPORTANT
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PICO 6.12: Risk of peri-procedural cranial nerve injury

6	randomised trials	not serious	not serious	serious ^a	not serious	very strong association ^a	7/1892 (0.4%)	103/1877 (5.5%)	RR 0.10 (0.05 to 0.20)	49 fewer per 1,000 (from 52 fewer to 44 fewer)	 HIGH	IMPORTANT
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CI: Confidence interval; RR: Risk ratio

Explanations

- Stenting and endarterectomy have evolved since the time of the contributing trials
- Few events, wide confidence intervals
- Significant heterogeneity, $I^2 > 60\%$
- Very wide confidence intervals
- Large effect

Table 7. Synoptic table of all recommendations

Recommendations	Quality of evidence	Strength of recommendation
In patients with $\geq 60\%$ asymptomatic carotid artery stenosis considered to be at increased risk of stroke on best medical therapy alone, we recommend carotid endarterectomy.	Moderate $\oplus\oplus\oplus$	Strong for carotid endarterectomy $\uparrow\uparrow$
In patients with asymptomatic carotid stenosis, recommend against carotid artery stenting as a routine alternative to best medical therapy alone.	Very low \oplus	Weak against carotid stenting \downarrow
In patients with asymptomatic carotid stenosis in whom revascularisation is considered to be appropriate, we suggest endarterectomy as the current treatment of choice.	Moderate $\oplus\oplus\oplus$	Weak for carotid endarterectomy \uparrow
In patients with severe (70-99%) symptomatic carotid artery stenosis, we recommend carotid endarterectomy.	Moderate $\oplus\oplus\oplus$	Strong for carotid endarterectomy $\uparrow\uparrow$
In patients with moderate (50-69%) symptomatic carotid artery stenosis, we suggest carotid endarterectomy.	Low $\oplus\oplus$	Weak for carotid endarterectomy \uparrow
In patients with mild ($<50\%$) symptomatic carotid artery stenosis, we recommend against carotid endarterectomy.	Very low \oplus	Strong against carotid endarterectomy $\downarrow\downarrow$
In patients with 50-99% symptomatic carotid stenosis in whom surgery is considered appropriate, we recommend early endarterectomy, ideally within two weeks of the last neurological event.	High $\oplus\oplus\oplus\oplus$	Strong for carotid endarterectomy $\uparrow\uparrow$
In patients with symptomatic carotid artery stenosis requiring revascularisation, we recommend endarterectomy as the treatment of choice.	Moderate $\oplus\oplus\oplus$	Strong for carotid endarterectomy $\uparrow\uparrow$
In patients with symptomatic carotid stenosis <70 years old requiring revascularisation, we suggest that stenting may be considered as an alternative to endarterectomy.	Low $\oplus\oplus$	Weak for carotid stenting \uparrow

Table 8. Synoptic table of all expert consensus statements

Expert consensus statements	Based on voting by all MWG members	Voting results
In selected patients 75 years of age or older with $\geq 60\%$ asymptomatic carotid artery stenosis and an expected survival of at least five years, who are considered to be at an increased risk of stroke on best medical therapy alone, carotid endarterectomy is suggested after careful consideration of the risks and benefits at a multi-disciplinary team meeting.		12/12
In patients with asymptomatic carotid stenosis in whom revascularisation is considered to be appropriate and who are less suitable for surgery, stenting may be suggested. We recommend careful consideration of the risks and benefits at a multi-disciplinary team meeting.		12/12
The independently assessed risk of in-hospital stroke or death following endarterectomy or stenting for asymptomatic carotid stenosis should be as low as possible, ideally below 2%.		12/12
The suitability of a patient with symptomatic carotid stenosis for carotid endarterectomy versus stenting should also take into account the interval since their last ischaemic cerebrovascular event, as well as anatomical and morphological features, including the atherosclerotic burden of the aortic arch.		12/12
The independently assessed risk of in-hospital stroke or death following endarterectomy or stenting for symptomatic carotid stenosis should be as low as possible, ideally below 4%.		11/12
Where possible, the indication for carotid endarterectomy or carotid artery stenting should be discussed at a multi-disciplinary team meeting. Consensus decisions can be made in between meetings, in order not to delay urgent revascularisations.		12/12
12/12 experts concluded that the establishment of validated local, regional or national registries, including audit systems for carotid interventions to monitor complication rates in patients with asymptomatic and symptomatic carotid stenosis is recommended.		12/12
MWG: Module Working Group		

Discussion

This evidence-based guideline was developed following the GRADE process and provides recommendations for the treatment of symptomatic and asymptomatic carotid stenosis by endarterectomy (CEA) or stenting (CAS) versus best medical therapy alone. All recommendations and expert consensus statements are summarised in table 7 and table 8.

Carotid revascularisation has been studied in randomised clinical trials for more than three decades, providing a wealth of evidence. Observational case series and large-scale registries are important to advance treatments and provide contemporary data on risks in real-world settings, but ultimately, the choice between treatment options should be informed by evidence from high quality RCTs, where such trials are available. We therefore based our recommendations in this guideline document for the choice between medical therapy alone, CEA or CAS on the evidence derived from randomised clinical trials only.

In some areas, particularly for stenting of asymptomatic carotid stenosis, the available evidence from clinical trials is still limited. However, additional data from large trials in asymptomatic carotid stenosis which are currently ongoing are expected in the near future and should provide a stronger evidence base to guide management of these patients.

CAS and CEA differ in treatment-associated risks, such as myocardial infarction and stroke. To fully determine the overall clinical impact of these outcomes in patients, additional measures such as quality of life and level of dependency should be systematically assessed in future trials.

We also acknowledge the fact that many of the trials providing the evidence for these guidelines were performed two to three decades ago. There have been important advances in the medical management of patients with atherosclerosis, and technical developments have also improved the safety of CEA and CAS since then. Because we had some concerns – especially in patients with asymptomatic carotid stenosis - that the applicability of the findings obtained in earlier trials may not apply to current clinical practice with contemporary medical and interventional treatment, we reduced the grade of some of the evidence for “indirectness”.

Any benefit of CEA or CAS is closely related to peri-procedural complication rates. Since the in-hospital complication rates of CEA and CAS have improved in recent years, expert consensus statements were

prepared which suggested that the independently-assessed peri-operative stroke and death rates after CEA or CAS should ideally be below 2% in patients with asymptomatic carotid stenosis and below 4% in patients symptomatic carotid stenosis. In randomised trials, about two thirds of these events occurred in the first two days after treatment, when patients were typically still in hospital.(92) Therefore these proposed acceptable in-hospital thresholds of 2% and 4% correspond with the traditionally-recommended 30-day thresholds of 3% and 6% for patients with asymptomatic and symptomatic stenosis, respectively. In-hospital thresholds may be more easily applicable to routine clinical practice because many patients will not be independently assessed by a neurologist or stroke physician 30 days after intervention. Moreover, outcomes following CEA and CAS should ideally be analysed at a local, regional and national level.

With modern medical management aiming for lower targets for lipid and blood pressure control, and more effective antiplatelet regimens (especially in patients with recent symptoms), the risk of stroke in asymptomatic and symptomatic carotid stenosis is expected to be lower than in the medical arms of some prior published trials. Ongoing trials are investigating whether contemporary medical therapy may obviate the need for invasive revascularisation in selected patient groups.

There have been a number of developments in the field of carotid artery stenting since the first trials which compared stenting with endarterectomy were completed, including the design of closed-cell and mesh-design stents,(93, 94) newer approaches to cerebral protection (involving reversal or arrest of blood flow),(95-106) and alternative access routes which avoid the aortic arch (including trans-brachial and trans-carotid access).(107-111) In addition, quality assurance programmes for stenting have been introduced in some countries.(112) For patients with symptomatic stenosis, the restriction to the evidence from past randomised trials may underestimate the role of CAS in experienced centres who are able to maintain low peri-procedural complication rates. Although stenting using more modern state-of-the-art techniques might reduce the peri-procedural risk of stroke, this needs to be tested in randomised trials of CAS versus CEA. Until further evidence is available, in patients requiring carotid revascularisation, the current weight of evidence is in favour of recommending CEA over CAS in most patient subgroups.

Plain language summary

Carotid stenosis refers to narrowing of a major blood vessel in the neck (the carotid artery) which carries blood to the eye and brain and is caused by fatty and calcium deposits in the blood vessel wall (atherosclerotic plaque). Carotid stenosis may cause a transient ischaemic attack (TIA or 'warning stroke') or a stroke. The narrowing can be removed by a surgical procedure called 'carotid endarterectomy', during which the surgeon opens the artery and removes the carotid plaque. An alternative treatment, called 'carotid artery stenting', involves passing a fine wire and tube through the skin and into the narrowed artery in the neck. A metal tube (stent) is placed inside the carotid artery to open it up with a view to preventing it from narrowing again. In patients who have not experienced recent symptoms (such as stroke, TIA, or ocular (eye) symptoms) from their carotid stenosis ('asymptomatic patients'), but who are still considered to be at risk of stroke on medication alone, we recommend carotid endarterectomy. In patients who have recently experienced these symptoms ('symptomatic patients'), we recommend carotid endarterectomy if the stenosis is severe, and suggest carotid endarterectomy may be considered if the stenosis is moderate. If surgery is recommended, we advise that carotid endarterectomy should be carried out as early as possible after the patient's initial symptoms, preferably within 2 weeks. Carotid artery stenting can be considered as an option to carotid endarterectomy in patients with symptomatic carotid stenosis, especially in patients younger than 70 years of age.

Acknowledgements

We would like to acknowledge Guillaume Turc (chair of the ESO guidelines board), Thorsten Steiner (former chair of the ESO guidelines board), Anne Hege Aamodt and Terry Quinn for reviewing the PICO's; and Bart van der Worp (past president of the ESO), Simona Sacco (the co-chair of the ESO guidelines board), Anne Hege Aamodt, Christian Nolte, Laura Capoccia and George Hamilton for reviewing the final text. We would also like to acknowledge Luzia Balmer for her excellent administrative support.

Figures

1. Endarterectomy or medical therapy for asymptomatic carotid stenosis

Figure 1.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death in endarterectomy versus medical therapy for asymptomatic carotid stenosis

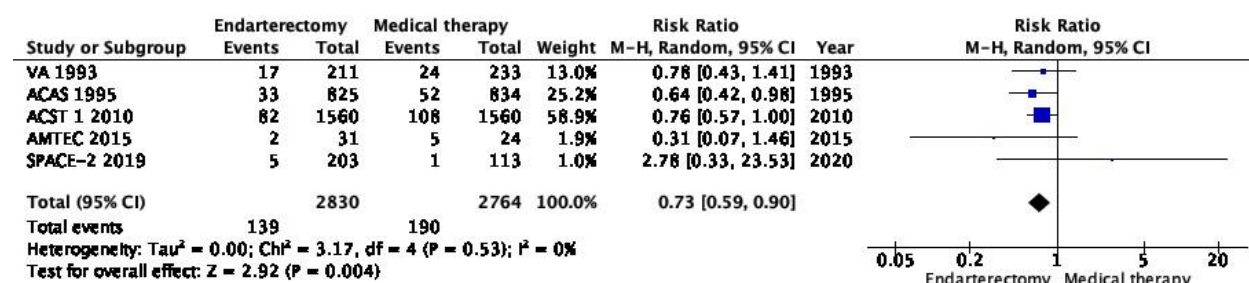
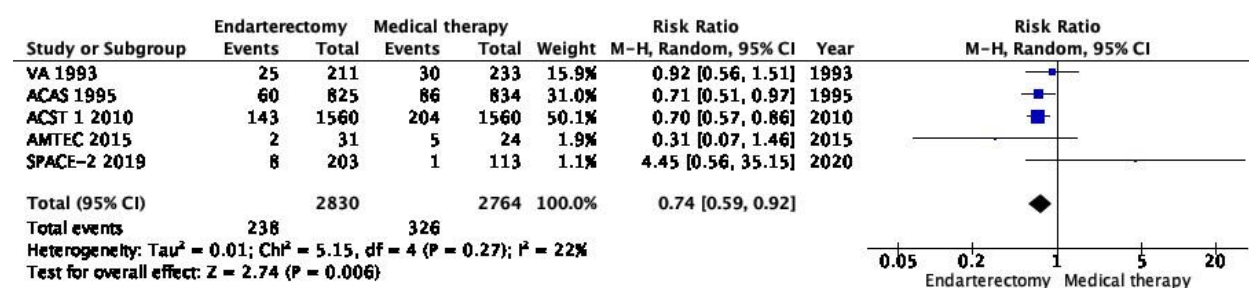


Figure 1.2: Long-term risk of stroke in any territory, including peri-procedural death in endarterectomy versus medical therapy for asymptomatic carotid stenosis



Subgroup analyses on PICO question 1.2: Long-term risk of stroke in any territory, including peri-procedural death in endarterectomy versus medical therapy for asymptomatic carotid stenosis.

Figure 1.2.1: Subgroup: Sex

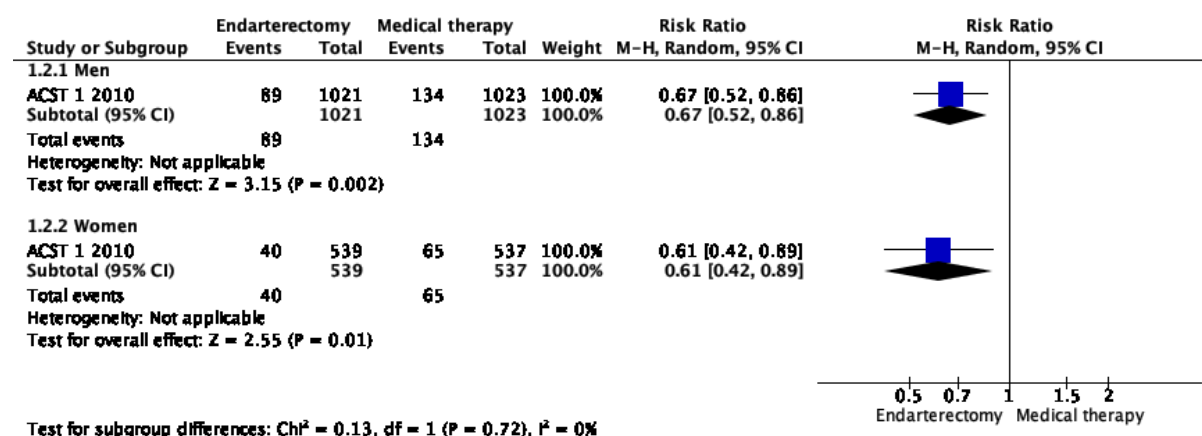


Figure 1.2.2: Subgroup: Age

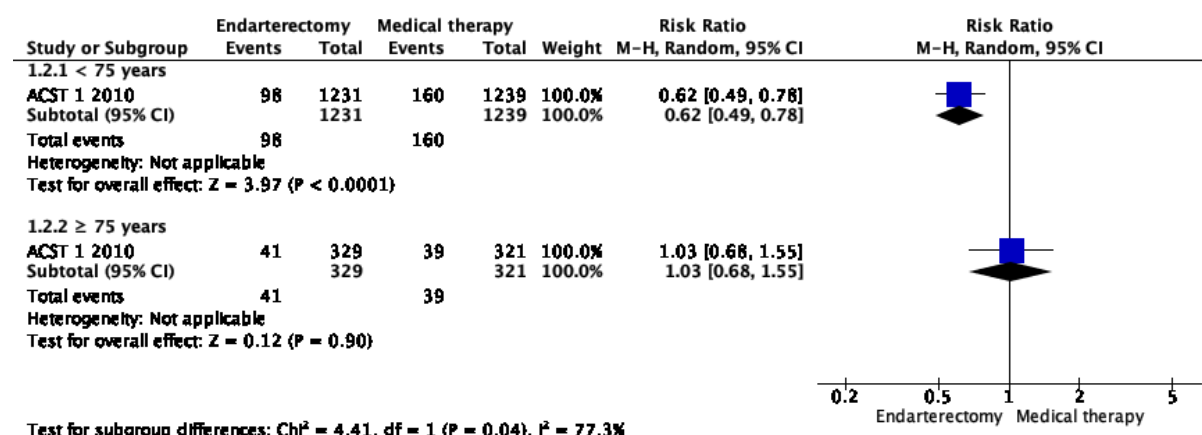


Figure 1.2.3: Subgroup: Severity of carotid stenosis

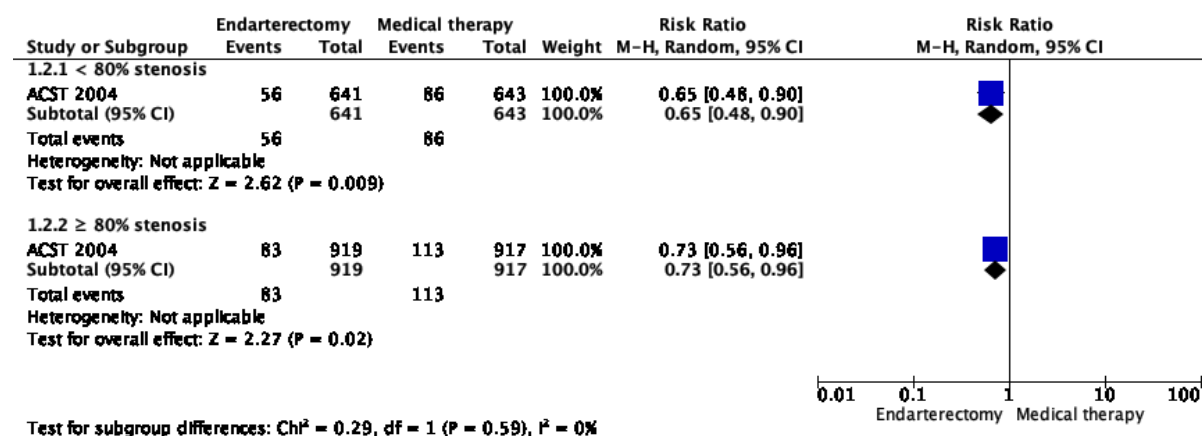


Figure 1.3: Long-term risk of major stroke, including peri-procedural death in endarterectomy versus medical therapy for asymptomatic carotid stenosis.

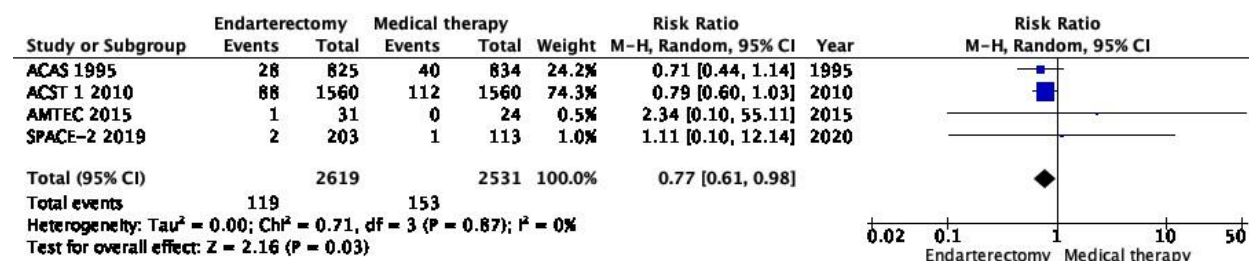
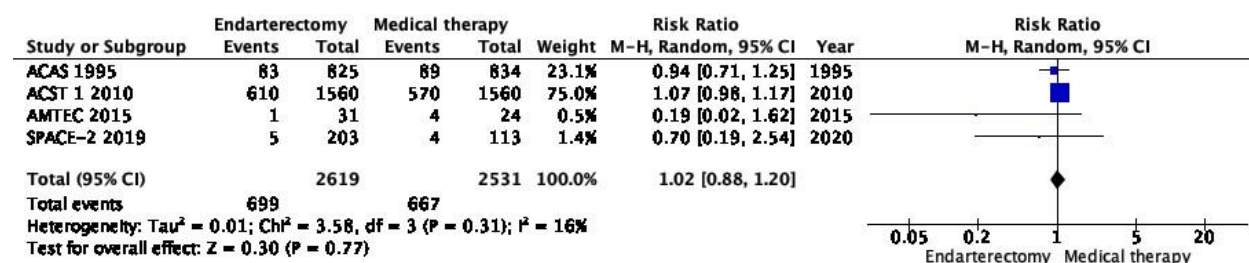


Figure 1.4: Long-term risk of death in asymptomatic in endarterectomy versus medical therapy for asymptomatic carotid stenosis



2. Stenting or medical therapy for asymptomatic carotid stenosis

Figure 2.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death in stenting versus medical therapy for asymptomatic carotid stenosis

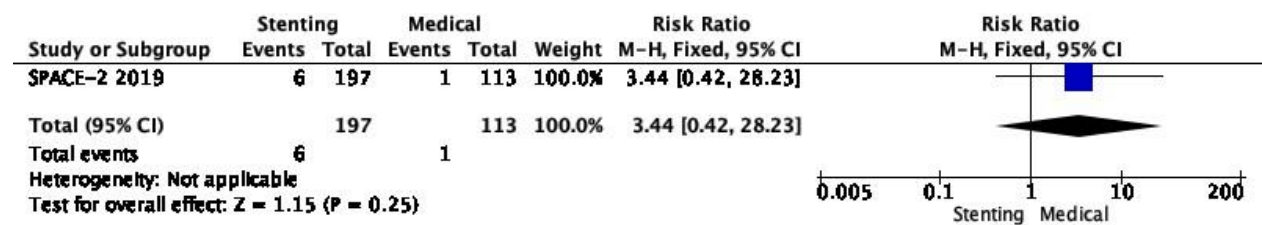


Figure 2.2: Long-term risk of stroke in any territory, including peri-procedural death including peri-procedural stroke or peri-procedural death in stenting versus medical therapy for asymptomatic carotid stenosis

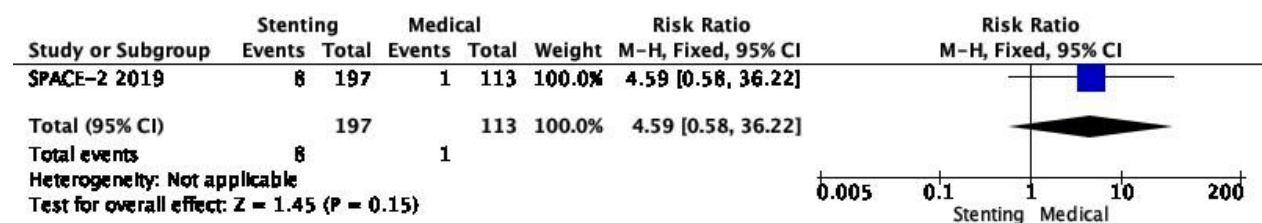


Figure 2.3: Long-term risk of major stroke, including peri-procedural death including peri-procedural stroke or peri-procedural death in stenting versus medical therapy for asymptomatic carotid stenosis

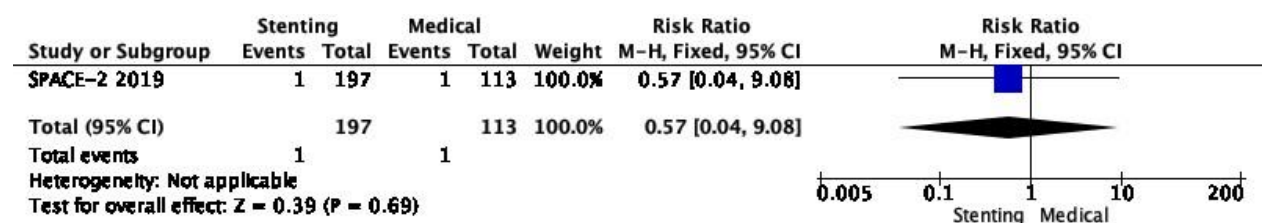
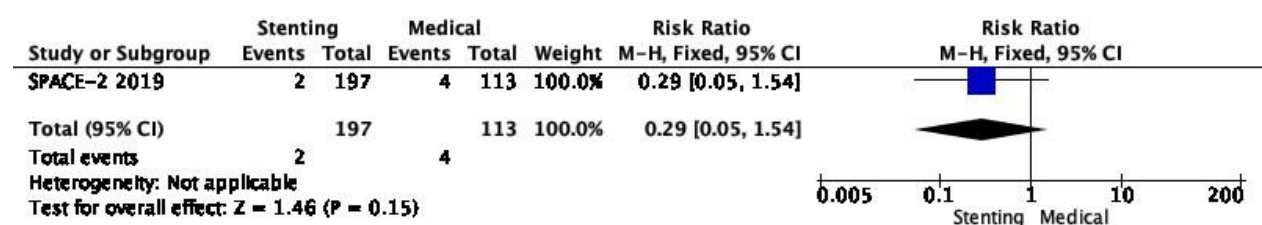


Figure 2.4: Long-term risk of death including peri-procedural stroke or peri-procedural death in stenting versus medical therapy for asymptomatic carotid stenosis



3. Stenting or endarterectomy for asymptomatic carotid stenosis

Figure 3.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death in stenting versus endarterectomy for asymptomatic carotid stenosis

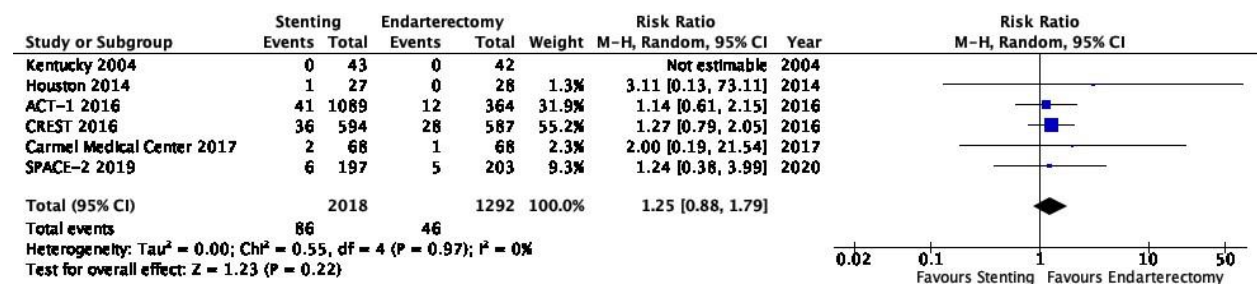


Figure 3.2: Long-term risk of post-procedural ipsilateral stroke in stenting versus endarterectomy for asymptomatic carotid stenosis

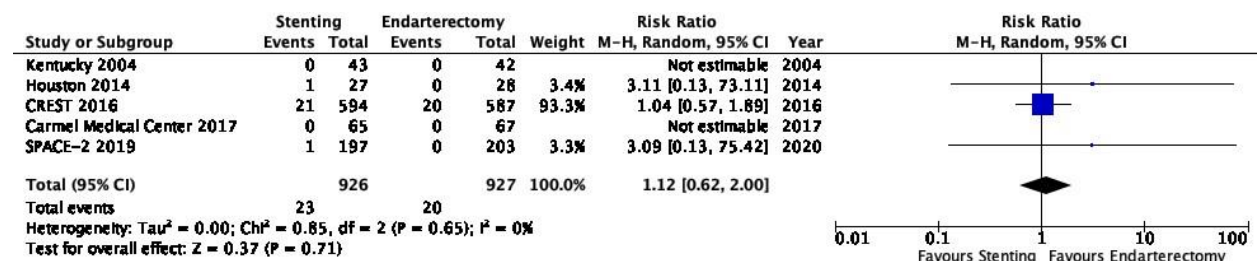


Figure 3.3: Long-term risk of stroke in any territory, including peri-procedural death in stenting versus endarterectomy for asymptomatic carotid stenosis

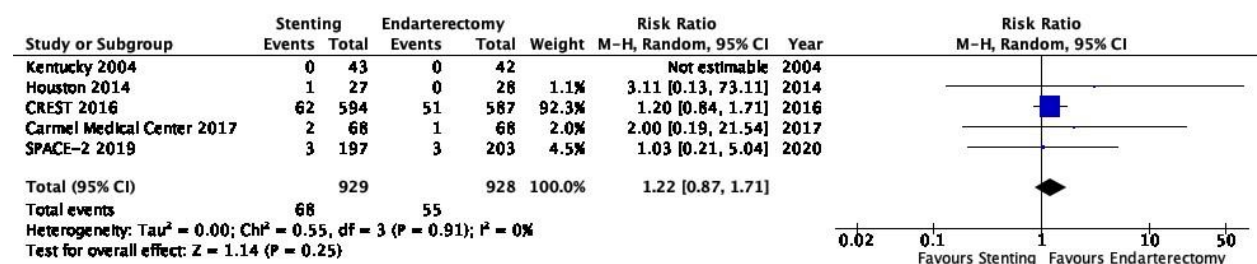


Figure 3.4: Long-term risk of major stroke, including peri-procedural death in stenting versus endarterectomy for asymptomatic carotid stenosis

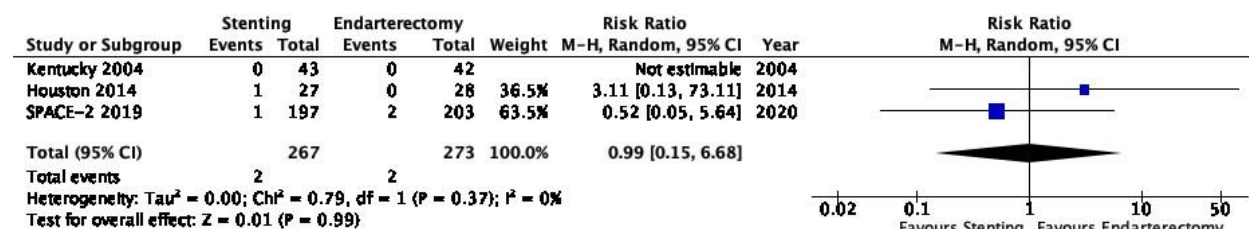


Figure 3.5: Long-term risk of death in stenting versus endarterectomy for asymptomatic carotid stenosis

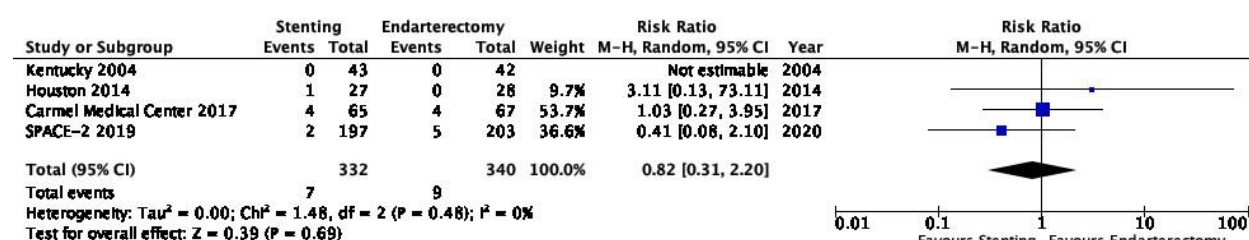


Figure 3.6: Peri-procedural stroke in stenting versus endarterectomy for asymptomatic carotid stenosis

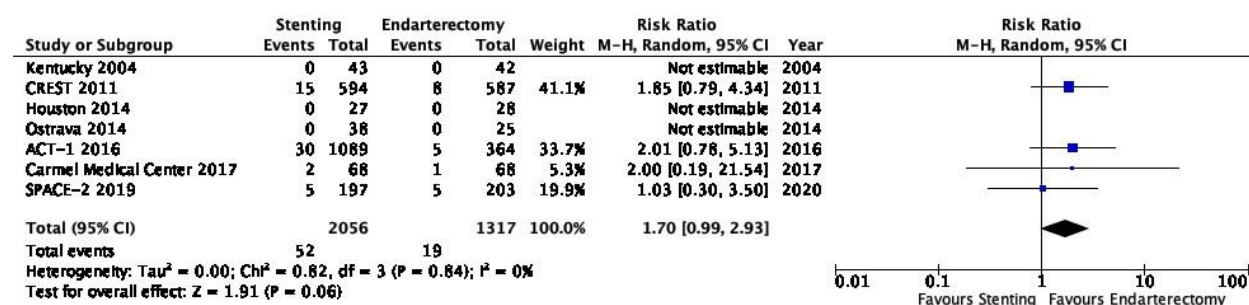


Figure 3.7: Peri-procedural death in stenting versus endarterectomy for asymptomatic carotid stenosis

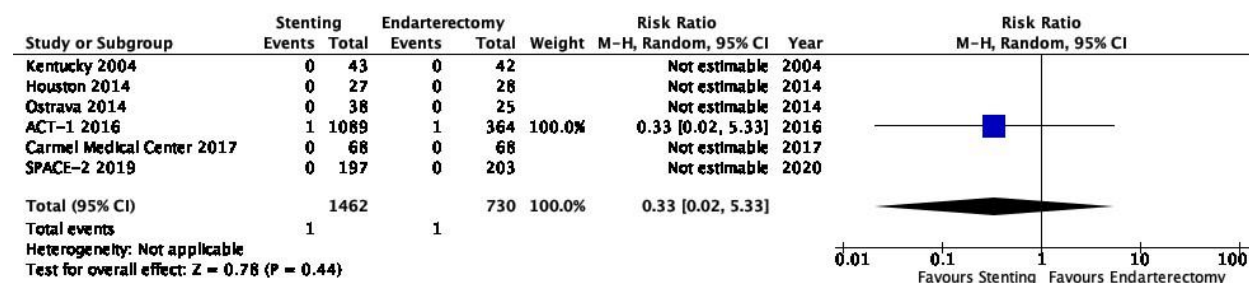


Figure 3.8: Peri-procedural stroke or death in stenting versus endarterectomy for asymptomatic carotid stenosis

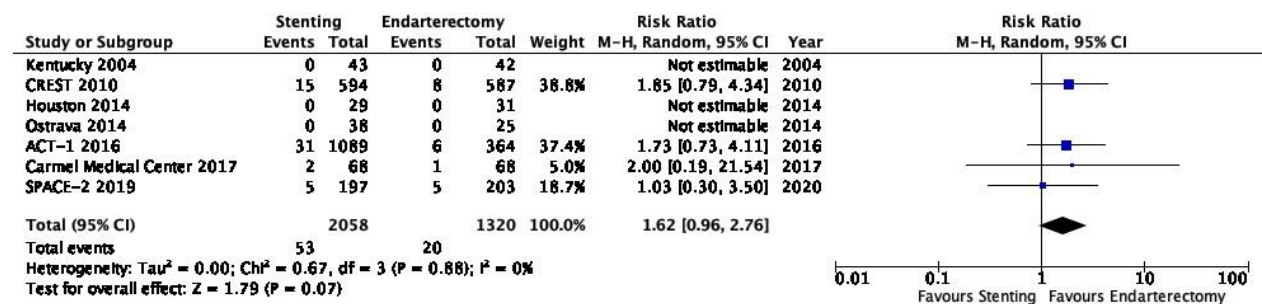


Figure 3.9: Peri-procedural major stroke or death in stenting versus endarterectomy for asymptomatic carotid stenosis

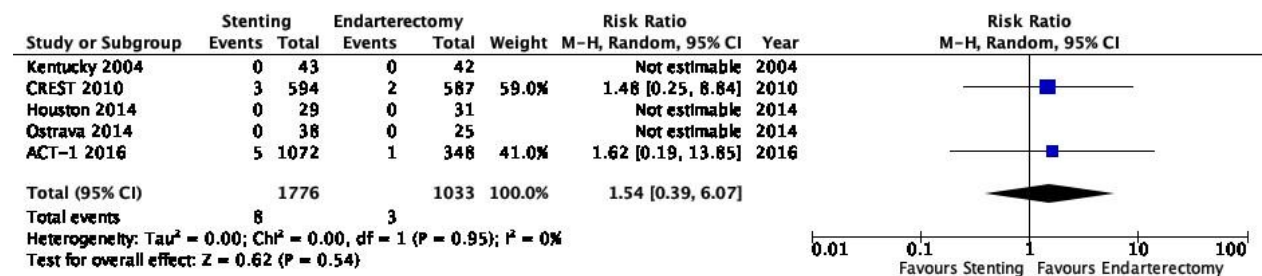


Figure 3.10: Peri-procedural myocardial infarction in stenting versus endarterectomy for asymptomatic carotid stenosis

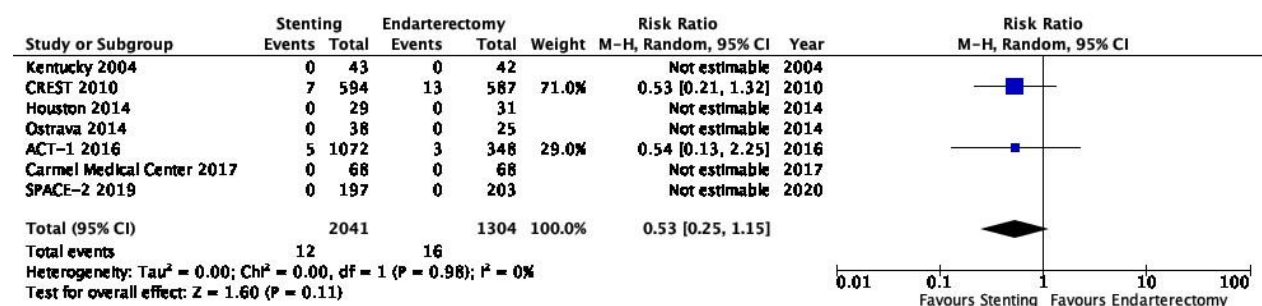
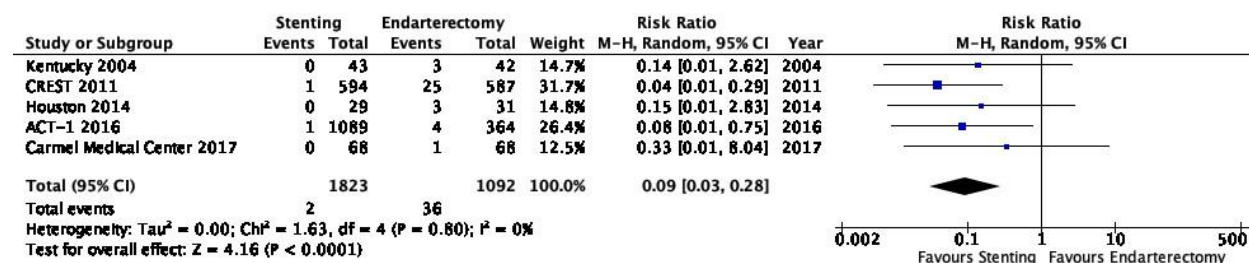
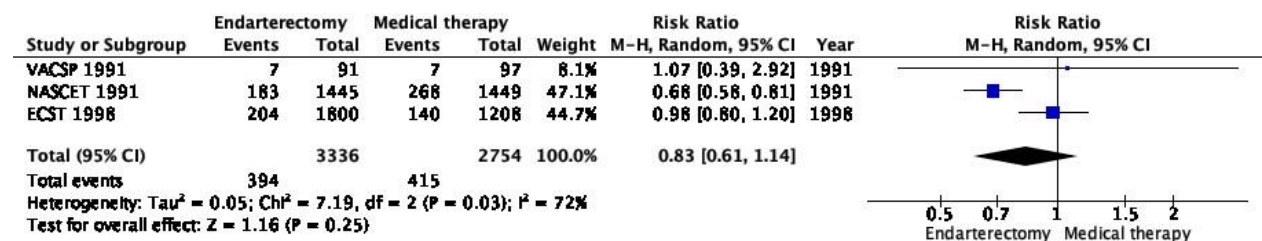


Figure 3.11: Peri-procedural cranial nerve injury in stenting versus endarterectomy for asymptomatic carotid stenosis



4. Endarterectomy or medical therapy for symptomatic carotid stenosis

Figure 4.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death in endarterectomy versus medical therapy for 30-99% symptomatic carotid stenosis



Subgroup analyses on PICO question 4.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke or peri-procedural death in endarterectomy versus medical therapy for 50-99% symptomatic carotid stenosis.

Figure 4.1.1: Subgroup: Age

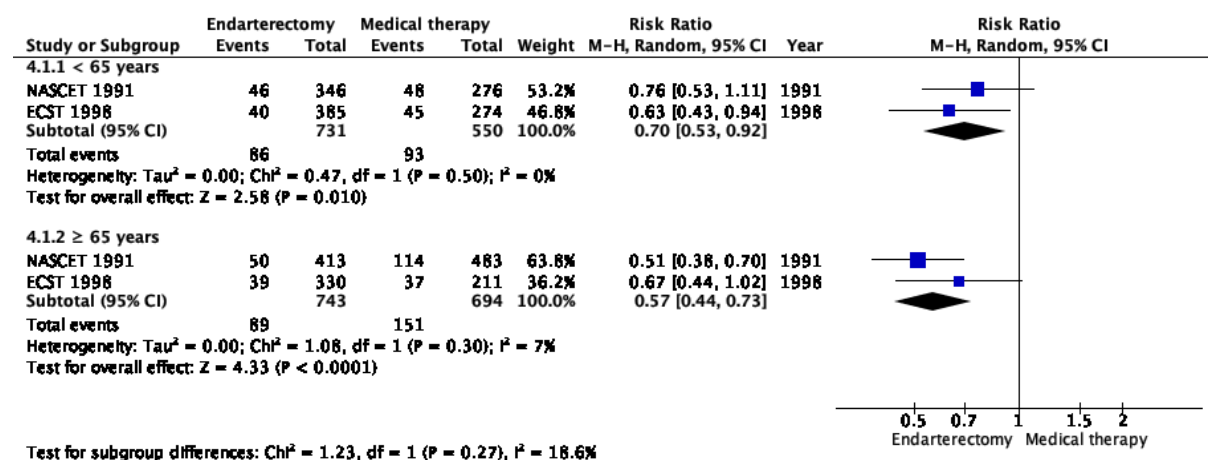


Figure 4.1.2: Subgroup: Sex

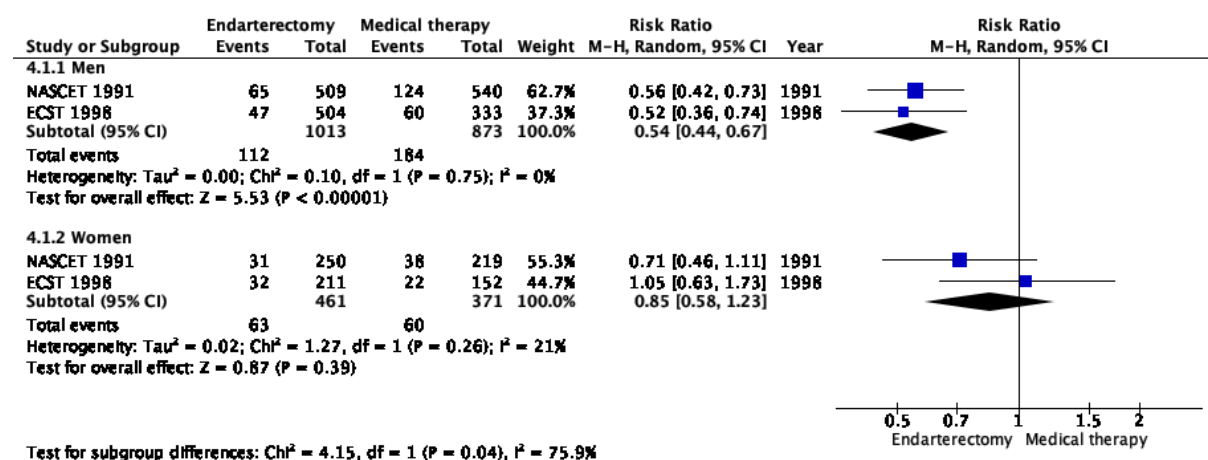


Figure 4.1.3: Subgroup: Time since last ischaemic event

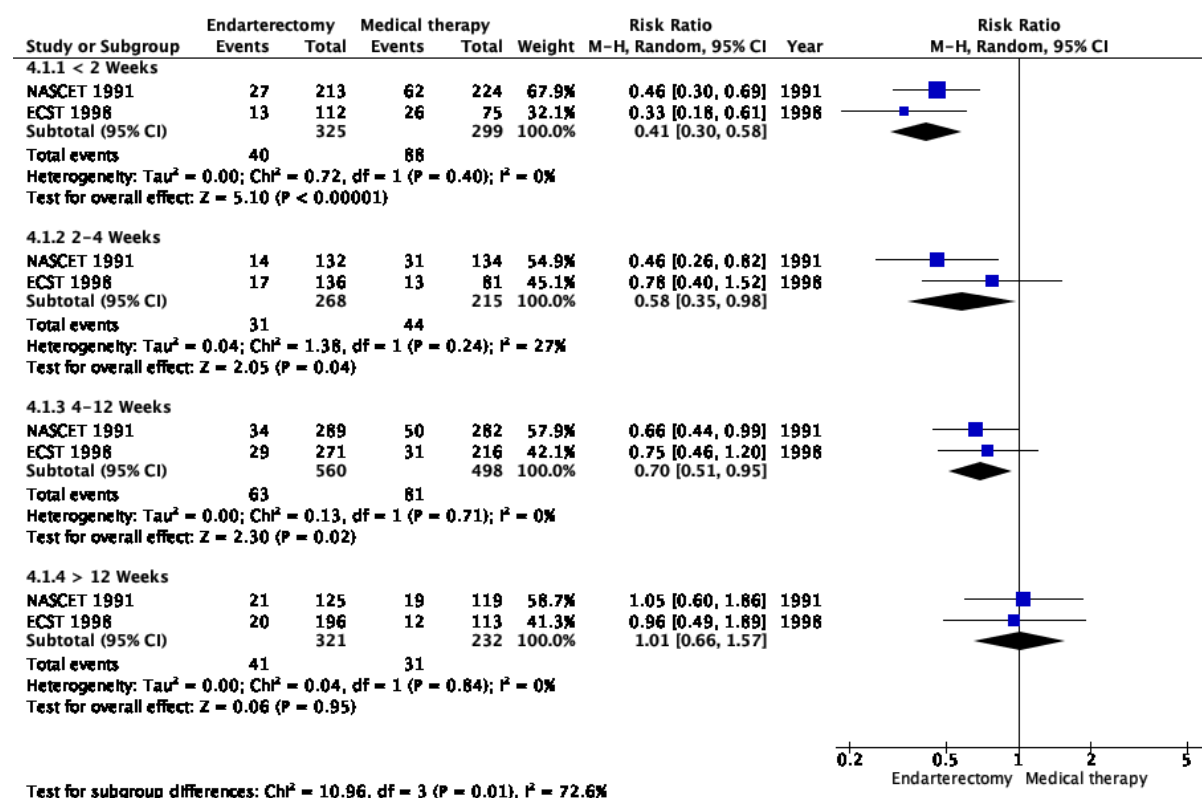


Figure 4.1.4: Subgroup: Severity of stenosis

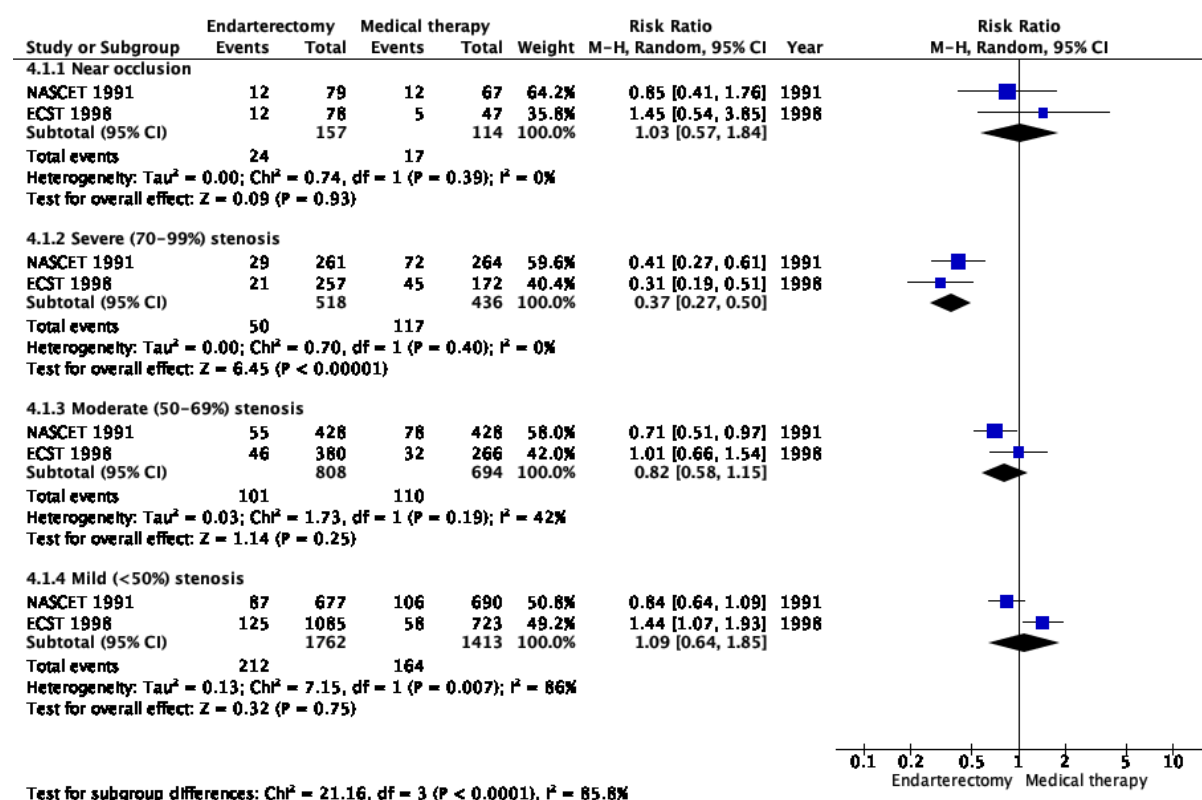
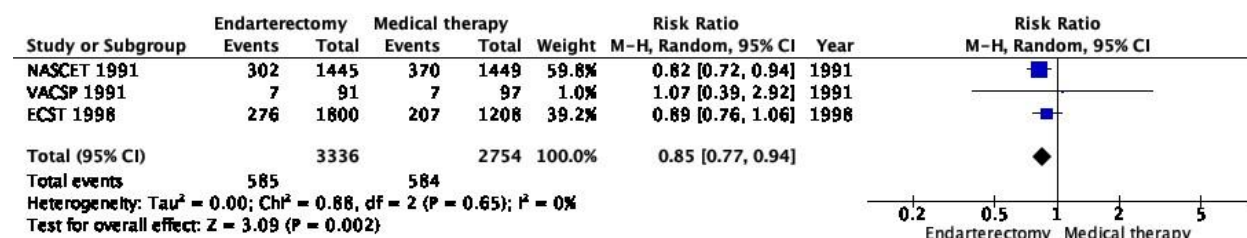


Figure 4.2: Long-term risk of stroke in any territory, including peri-procedural death in endarterectomy versus medical therapy for 30-99% symptomatic carotid stenosis



Subgroup analyses on PICO question 4.2: Long-term risk of stroke in any territory, including peri-procedural death in endarterectomy versus medical therapy for symptomatic carotid stenosis.

Figure 4.2.1: Subgroup: Severity of stenosis

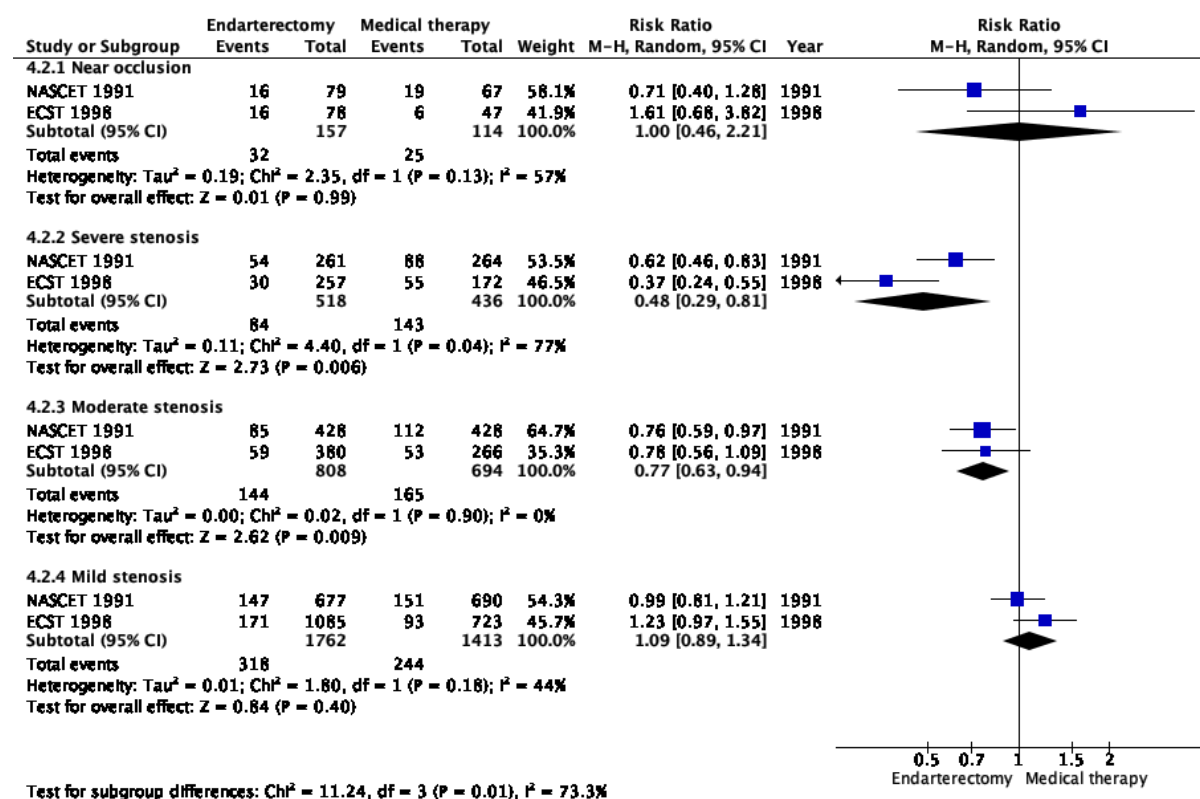
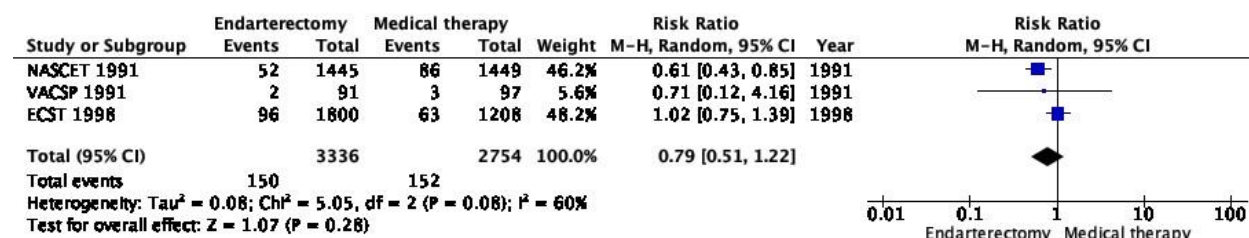


Figure 4.3: Long-term risk of major stroke, including peri-procedural death in endarterectomy versus medical therapy for 30-99% symptomatic carotid stenosis



Subgroup analysis on PICO question 4.3: Long-term risk of major stroke, including peri-procedural death in endarterectomy versus medical therapy for symptomatic carotid stenosis:

Figure 4.3.1: Subgroup: Severity of Stenosis

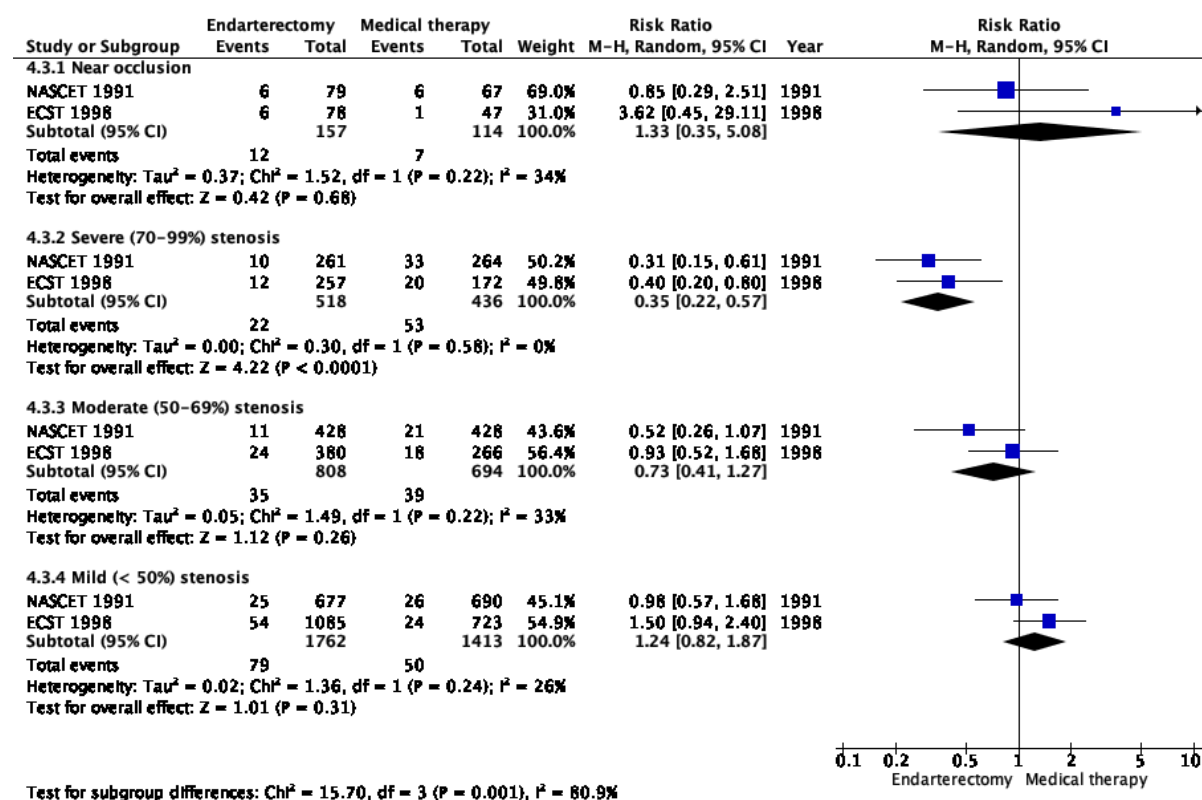
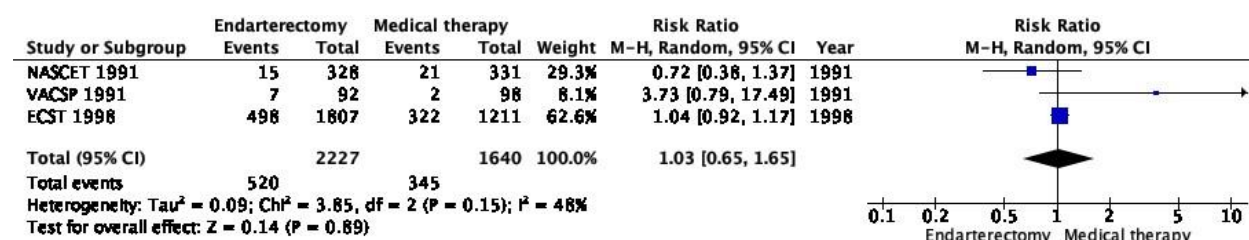


Figure 4.4: Long-term risk of death in endarterectomy versus medical therapy for 30-99% symptomatic carotid stenosis



5. Stenting or medical therapy for symptomatic carotid stenosis

No data available

6. Stenting or endarterectomy for symptomatic carotid stenosis

Figure 6.1: Long-term risk of ipsilateral stroke, including peri-procedural stroke in any territory or peri-procedural death in stenting versus endarterectomy for symptomatic carotid stenosis

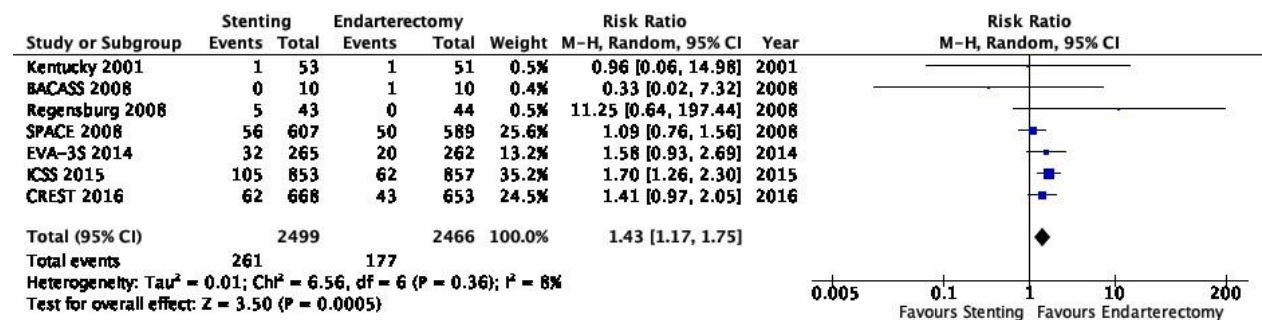


Figure 6.2: Long-term risk of post-procedural ipsilateral stroke in stenting versus endarterectomy for symptomatic carotid stenosis

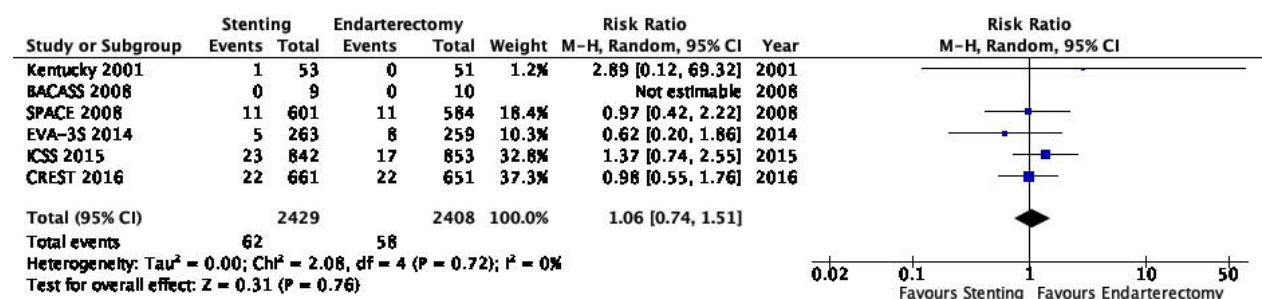


Figure 6.3: Long-term risk of stroke in any territory, including peri-procedural death in stenting versus endarterectomy for symptomatic carotid stenosis

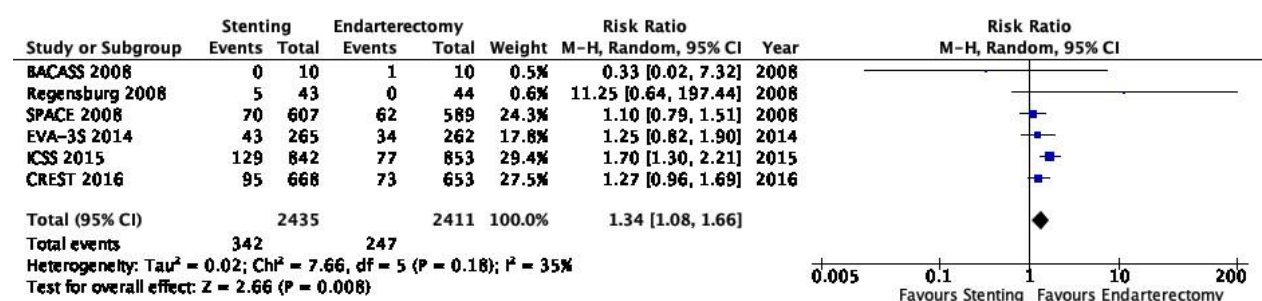


Figure 6.4: Long-term risk of major stroke, including peri-procedural death in stenting versus endarterectomy for symptomatic carotid stenosis

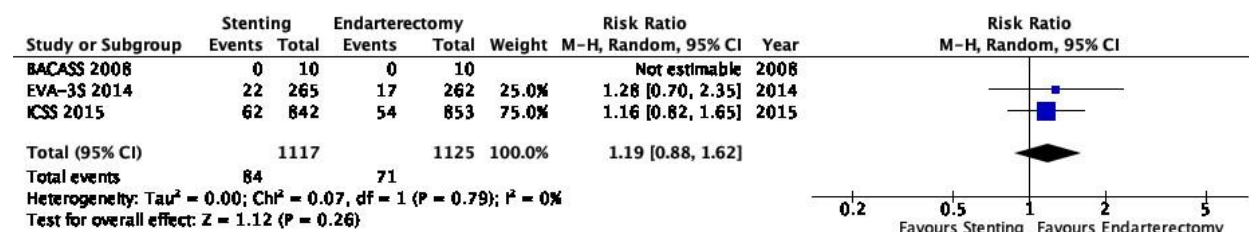


Figure 6.5: Long-term risk of death in stenting versus endarterectomy for symptomatic carotid stenosis

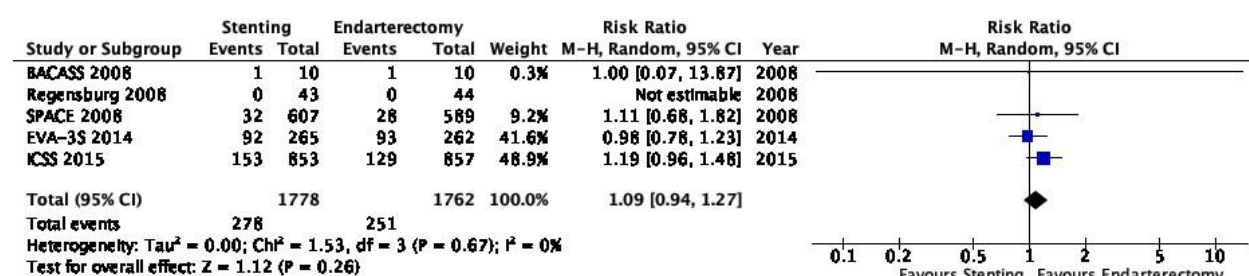


Figure 6.6: Long-term risk of severe restenosis in stenting versus endarterectomy for symptomatic or asymptomatic carotid stenosis

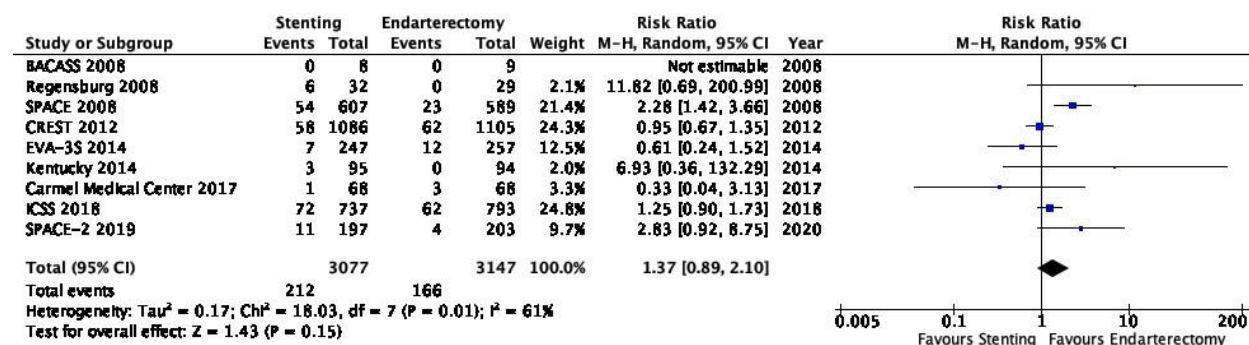


Figure 6.7: Peri-procedural stroke in stenting versus endarterectomy for symptomatic carotid stenosis

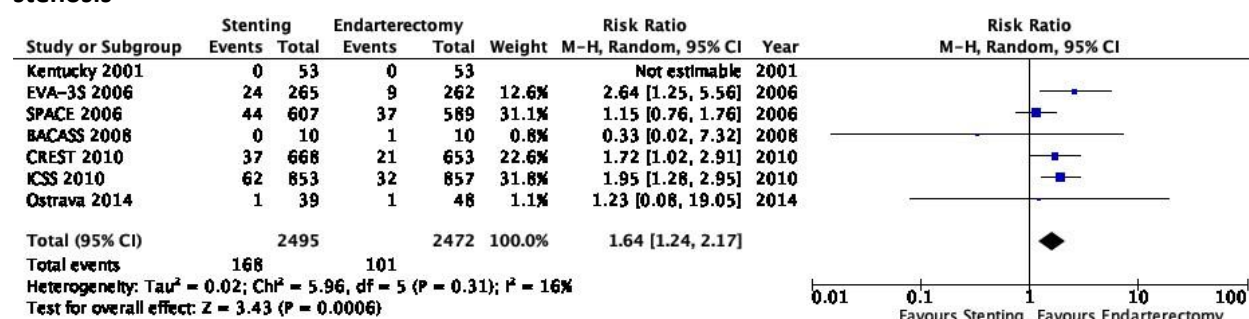


Figure 6.8: Peri-procedural death in stenting versus endarterectomy for symptomatic carotid stenosis

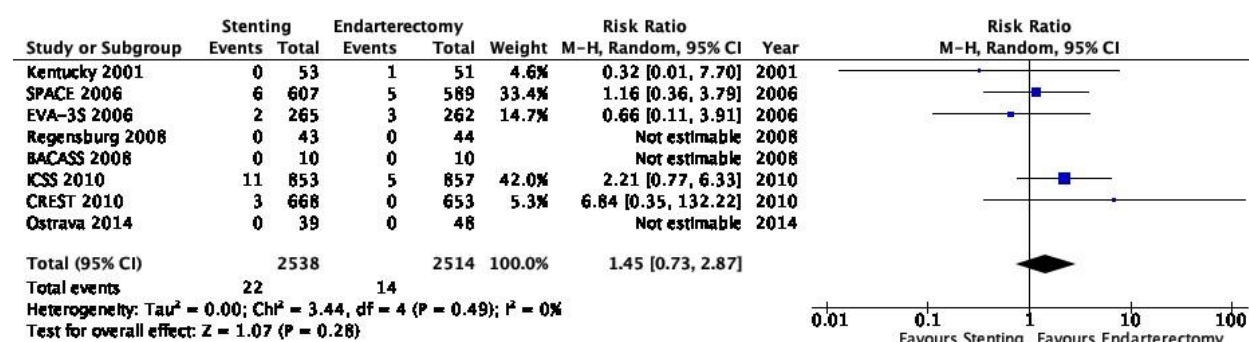
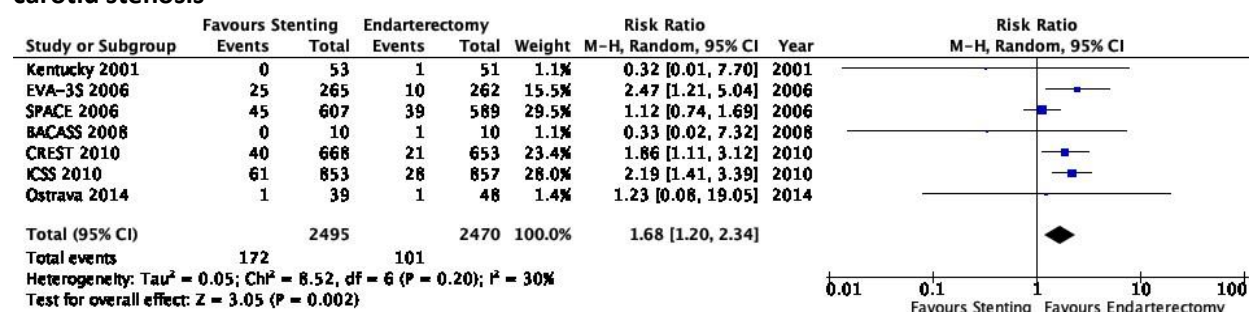


Figure 6.9: Peri-procedural stroke or death in stenting versus endarterectomy for symptomatic carotid stenosis



Subgroup analyses on PICO question 6.9: Peri-procedural stroke or death in stenting versus endarterectomy for symptomatic carotid stenosis.

Figure 6.9.1: Subgroup: Age

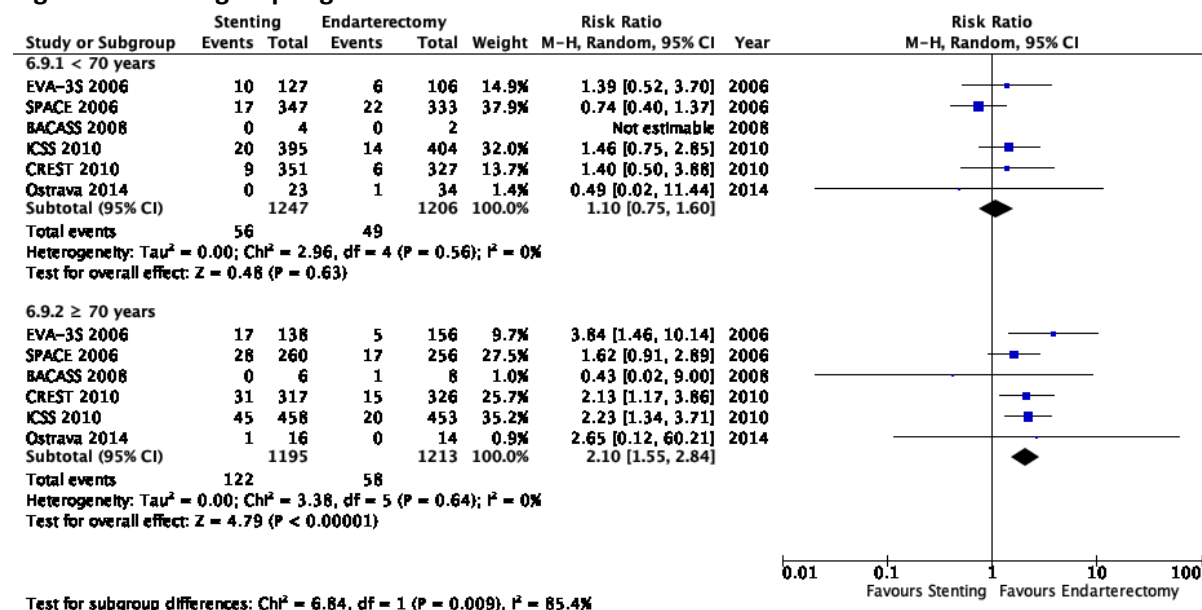


Figure 6.9.2: Subgroup: Sex

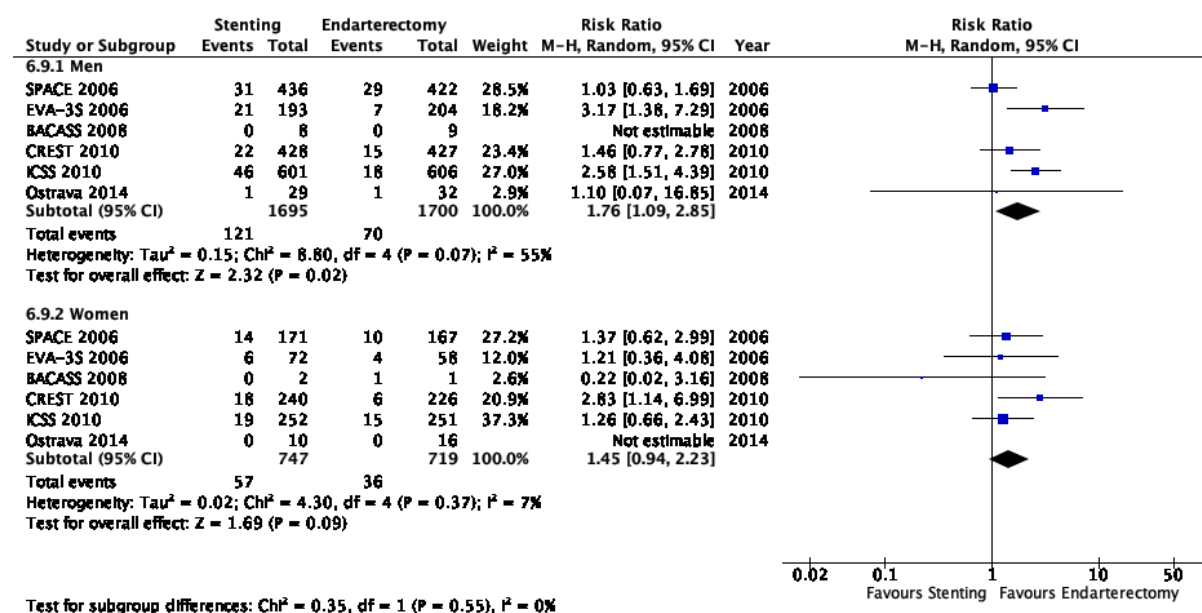


Figure 6.9.3: Subgroup: Severity of stenosis.

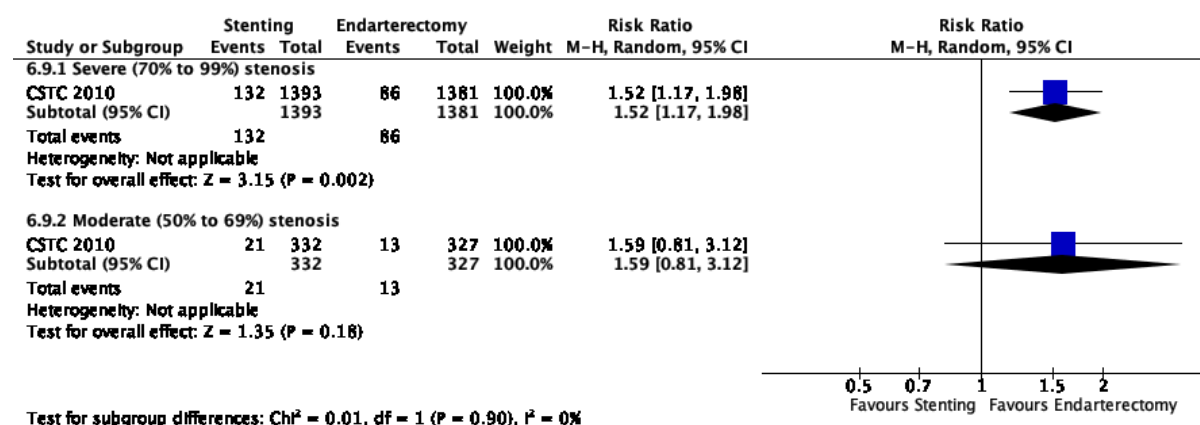


Figure 6.9.4: Subgroup: Time since last ischaemic event.

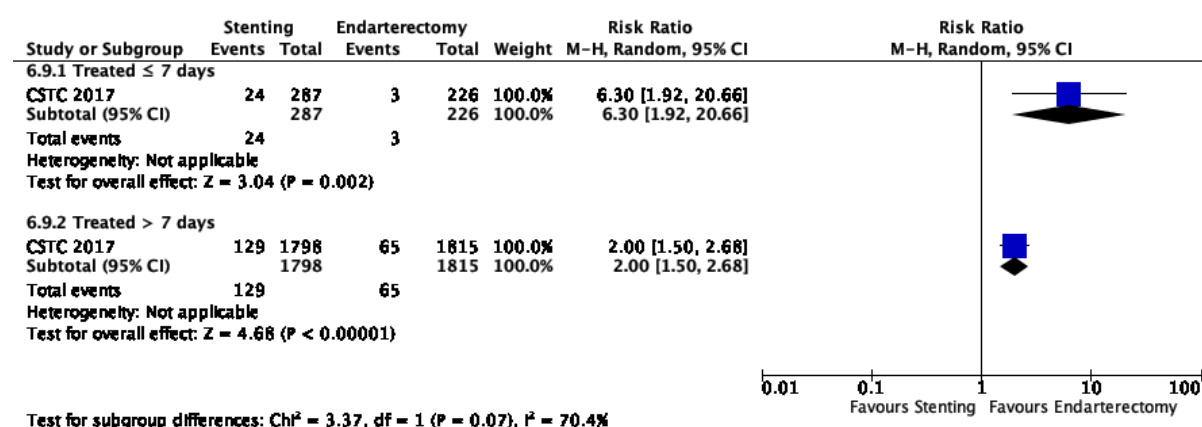


Figure 6.9.5: Subgroup: Type of last ischaemic event.

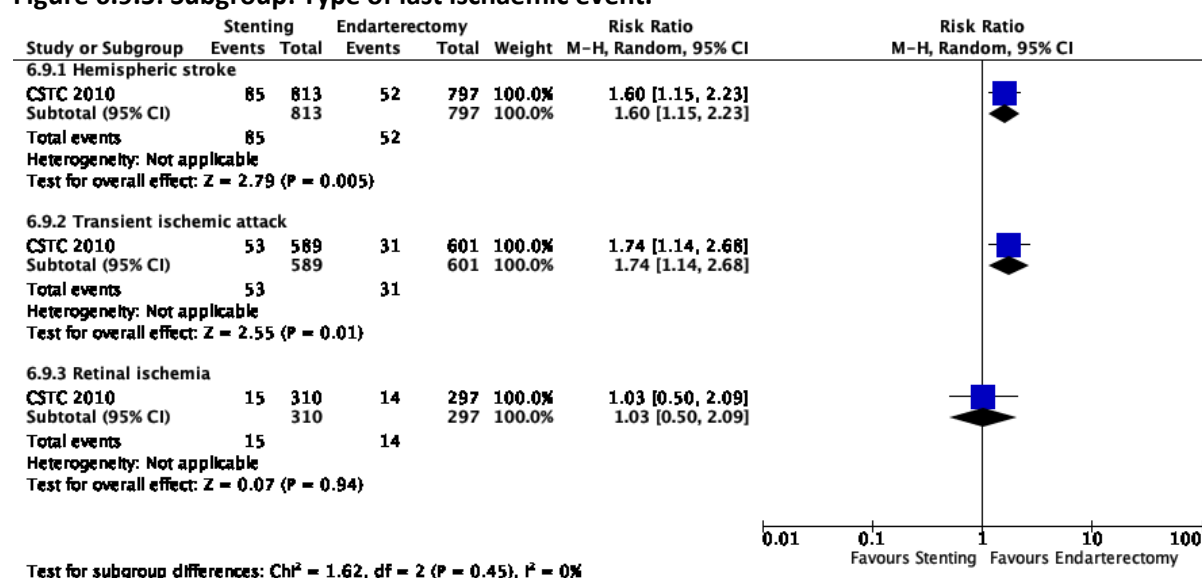


Figure 6.10: Peri-procedural major stroke or death in stenting versus endarterectomy for symptomatic carotid stenosis.

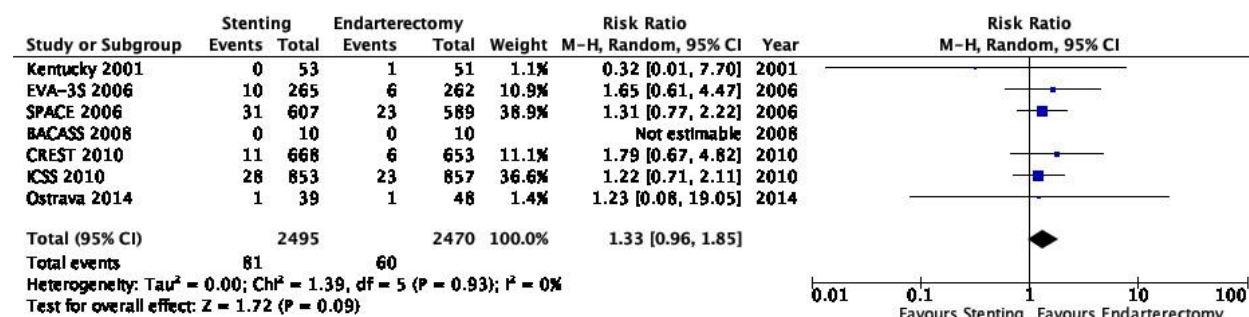


Figure 6.11: Peri-procedural myocardial infarction in stenting versus endarterectomy for symptomatic carotid stenosis.

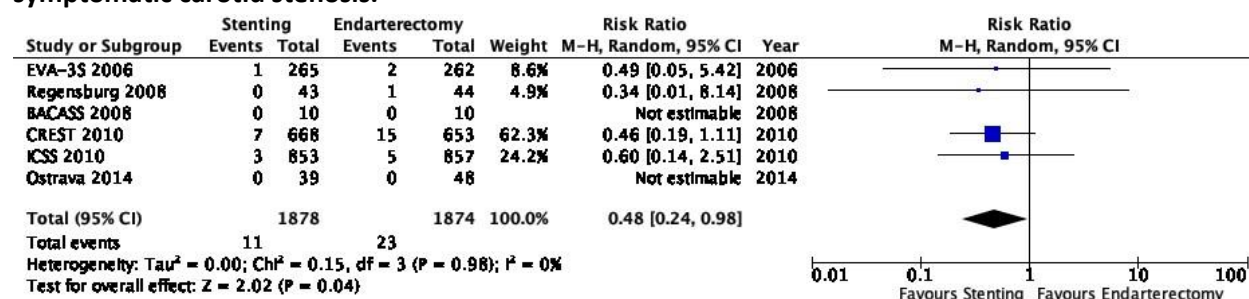
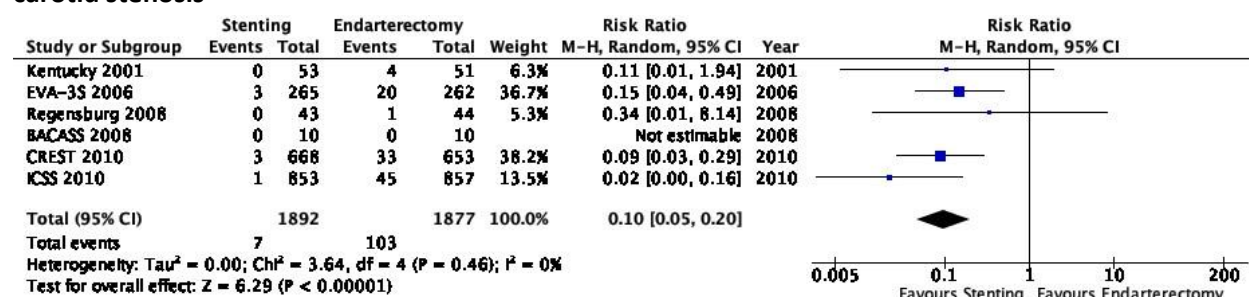


Figure 6.12: Peri-procedural cranial nerve injury in stenting versus endarterectomy for symptomatic carotid stenosis



Supplement

Literature search strategy

PICO 1: In patients with asymptomatic carotid stenosis, endarterectomy compared with medical therapy alone:

1. carotid stenosis/
2. carotid artery diseases/
3. carotid arteries/
4. carotid artery, common/
5. carotid artery, internal/
6. (carotid adj (arter\$ or stenosis)).tw
7. 1 or 2 or 3 or 4 or 5 or 6
8. endarterectomy, carotid/
9. endarterectomy/
10. (carotid adj5 endarterectomy).tw
11. 8 or 9 or 10
12. randomized controlled trials/
13. controlled clinical trials/
14. clinical trials/
15. random\$.tw
16. ((clin\$ or controlled) adj5 trial\$).tw
17. 12 or 13 or 14 or 15 or 16
18. 7 and 11 and 17
19. asymptomatic.tw
20. 18 and 19
- 21: Limit 20 to humans

PICO 2, 5: In patients with asymptomatic and symptomatic carotid stenosis, stenting compared with medical therapy

1. (carotid adj10 (stent\$ or angioplasty)).tw.
2. carotid artery diseases/ or carotid artery thrombosis/ or carotid stenosis/
3. carotid arteries/ or carotid artery, common/ or carotid artery, external/ or carotid artery, internal/
4. constriction, pathologic/
5. 3 and 4
6. (carotid adj5 (stenosis or thrombo\$ or disease\$ or narrow\$ or plaque\$ or arterioscler\$ or atheroscler\$)).tw.
7. 2 or 5 or 6
8. angioplasty/ or angioplasty, balloon/ or angioplasty, balloon, laser-assisted/ or angioplasty, laser/
9. Stents/
10. (angioplasty or stent\$ or endovascular).tw.
11. (balloon adj5 (dilat\$ or catheter\$)).tw.

12. ((endoluminal or transluminal) adj5 repair\$).tw.
13. 8 or 9 or 10 or 11 or 12
14. 7 and 13
15. 1 or 14
16. Randomized Controlled Trials as Topic/
17. random allocation/
18. Controlled Clinical Trials as Topic/
19. control groups/
20. clinical trials as topic/ or clinical trials, phase i as topic/ or clinical trials, phase ii as topic/ or clinical trials, phase iii as topic/ or clinical trials, phase iv as topic/
21. double-blind method/
22. single-blind method/
23. Placebos/
24. placebo effect/
25. Drug Evaluation/
26. Research Design/
27. randomized controlled trial.pt.
28. controlled clinical trial.pt.
29. clinical trial.pt.
30. random\$.tw.
31. (controlled adj5 (trial\$ or stud\$)).tw.
32. (clinical\$ adj5 trial\$).tw.
33. ((control or treatment or experiment\$ or intervention or surgical) adj5 (group\$ or subject\$ or patient\$)).tw.
34. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
35. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.
36. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
37. (coin adj5 (flip or flipped or toss\$)).tw.
38. latin square.tw.
39. versus.tw.
40. controls.tw.
41. or/16-40
42. 15 and 41
43. limit 42 to humans

PICO 4: In patients with symptomatic carotid stenosis, endarterectomy compared with medical therapy:

1. cerebrovascular disorders/ or basal ganglia cerebrovascular disease/ or brain ischemia/ or exp brain infarction/ or hypoxia-ischemia, brain/ or ischemic attack, transient/ or carotid artery diseases/ or carotid artery thrombosis/ or carotid artery, internal, dissection/ or carotid stenosis/ or exp carotid artery injuries/ or intracranial arterial diseases/ or cerebral arterial diseases/ or infarction, anterior cerebral artery/ or infarction, middle cerebral artery/ or infarction, posterior

cerebral artery/ or exp "intracranial embolism and thrombosis"/ or exp stroke/ or vertebral artery dissection/

2. (isch?emi\$ adj6 (stroke\$ or apoplex\$ or cerebral vasc\$ or cerebrovasc\$ or cva or attack\$)).tw.

3. ((brain or cerebr\$ or cerebell\$ or vertebrobasil\$ or hemispher\$ or intracran\$ or intracerebral or infratentorial or supratentorial or middle cerebr\$ or mca\$ or anterior circulation) adj5 (isch?emi\$ or infarct\$ or thrombo\$ or emboli\$)).tw.

4. (transient isch\$ or TIA or TIAs).tw.

5. 1 or 2 or 3 or 4

6. (carotid adj5 (stenosis or thrombo\$ or disease\$ or arter\$ or atherosclero\$ or atheroma\$ or narrow\$ or plaque\$ or occlus\$ or occlud\$ or constrict\$ or emboli* or block\$)).tw.

7. carotid artery diseases/su or carotid artery thrombosis/su or carotid artery, internal, dissection/su or carotid stenosis/su or exp carotid artery injuries/su or exp Carotid Arteries/su

8. carotid artery diseases/ or carotid artery thrombosis/ or carotid artery, internal, dissection/ or carotid stenosis/ or exp carotid artery injuries/ or exp Carotid Arteries/

9. carotid.tw.

10. 6 or 7 or 8 or 9

11. CEA.tw.

12. Endarterectomy/

13. 11 or 12

14. 10 and 13

15. Endarterectomy, Carotid/

16. (carotid adj5 (endarterectomy or thromboendarterectomy or surgery or revasculari\$ or eversion)).tw.

16. 14 or 15 or 16

17. Randomized Controlled Trials as Topic/

18. random allocation/

19. Controlled Clinical Trials as Topic/

20. control groups/

21. clinical trials as topic/

22. randomized controlled trial.pt.

23. controlled clinical trial.pt.

24. clinical trial.pt.

25. (random\$ or RCT or RCTs).tw.

26. (controlled adj5 (trial\$ or stud\$)).tw.

27. (clinical\$ adj5 trial\$).tw.

28. ((control or treatment or experiment\$ or intervention) adj5 (group\$ or subject\$ or patient\$)).tw.

29. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.

30. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw. 31. trial.ti.

32. (assign\$ or allocat\$).tw.

33. controls.tw.

34. or/17-33

35. 5 and 16 and 34

36. Limit 35 to humans

PICO 3 and 6: In patients with asymptomatic and symptomatic carotid stenosis, do endarterectomy and stenting differ

1. (carotid adj10 (stent\$ or angioplasty)).tw.
2. carotid artery diseases/ or carotid artery thrombosis/ or carotid stenosis/
3. carotid arteries/ or carotid artery, common/ or carotid artery, external/ or carotid artery, internal/
4. constriction, pathologic/
5. 3 and 4
6. (carotid adj5 (stenosis or thrombo\$ or disease\$ or narrow\$ or plaque\$ or arterioscler\$ or atheroscler\$)).tw.
7. 2 or 5 or 6
8. angioplasty/ or angioplasty, balloon/ or angioplasty, balloon, laser-assisted/ or angioplasty, laser/ or endarterectomy/
9. Stents/
10. (angioplasty or stent\$ or endovascular or CEA or endarterectomy).tw.
11. (balloon adj5 (dilat\$ or catheter\$)).tw.
12. ((endoluminal or transluminal) adj5 repair\$).tw.
13. 8 or 9 or 10 or 11 or 12
14. 7 and 13
15. 1 or 14
16. Randomized Controlled Trials as Topic/
17. random allocation/
18. Controlled Clinical Trials as Topic/
19. control groups/
20. clinical trials as topic/ or clinical trials, phase i as topic/ or clinical trials, phase ii as topic/ or clinical trials, phase iii as topic/ or clinical trials, phase iv as topic/
21. double-blind method/
22. single-blind method/
23. Placebos/
24. placebo effect/
25. Research Design/
26. randomized controlled trial.pt.
27. controlled clinical trial.pt.
28. clinical trial.pt.
29. random\$.tw.
30. (controlled adj5 (trial\$ or stud\$)).tw.
31. (clinical\$ adj5 trial\$).tw.
32. ((control or treatment or experiment\$ or intervention or surgical) adj5 (group\$ or subject\$ or patient\$)).tw.
33. (quasi-random\$ or quasi random\$ or pseudo-random\$ or pseudo random\$).tw.
34. ((control or experiment\$ or conservative) adj5 (treatment or therapy or procedure or manage\$)).tw.

- 35. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj5 (blind\$ or mask\$)).tw.
- 36. (coin adj5 (flip or flipped or toss\$)).tw.
- 37. latin square.tw.
- 38. versus.tw.
- 39. controls.tw.
- 40. or/16-39
- 41. 15 and 40
- 42. limit 41 to humans

Supplementary figures

Figure S1: Literature search endarterectomy versus medical therapy

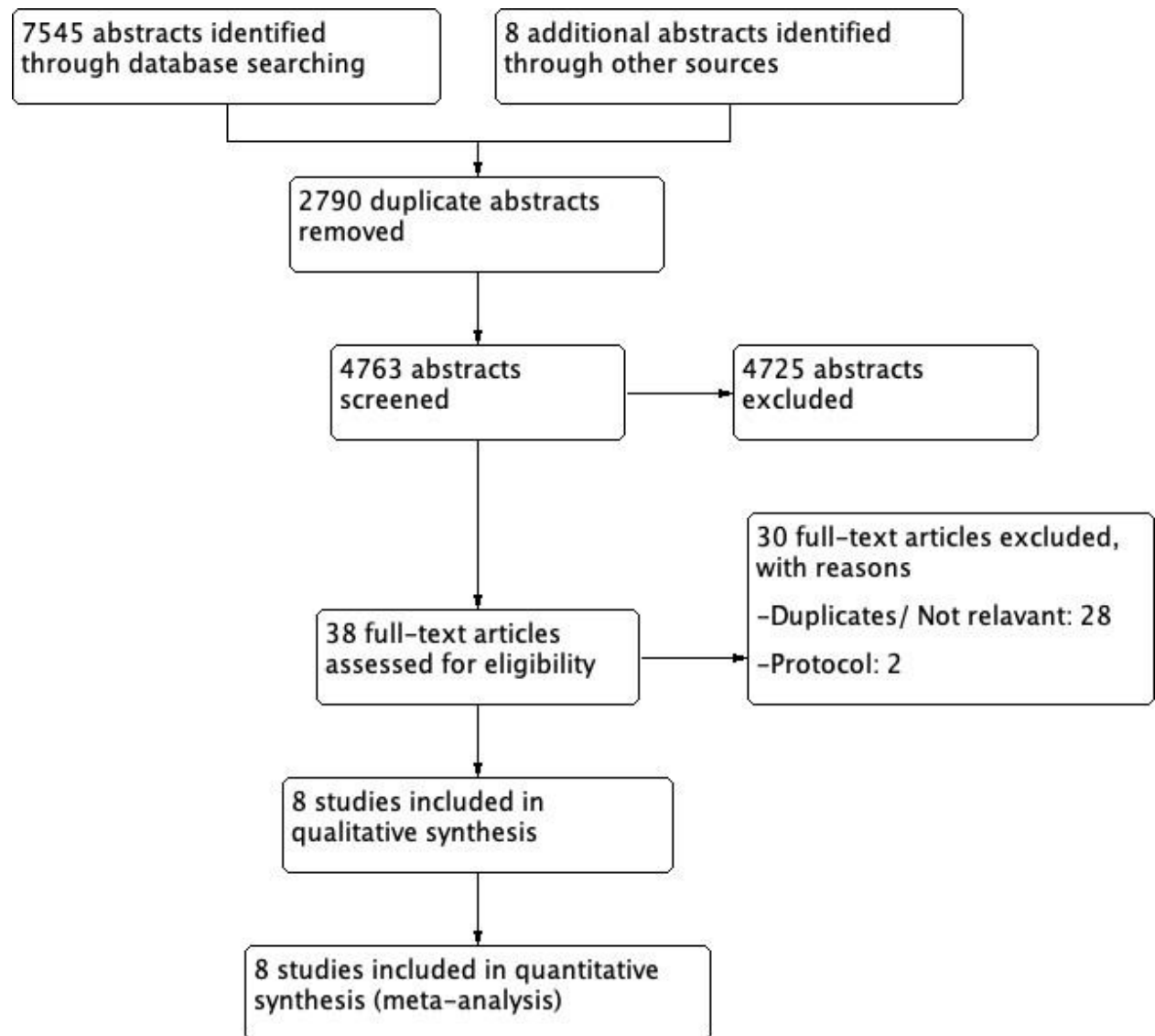


Figure S2: Literature search stenting versus medical therapy

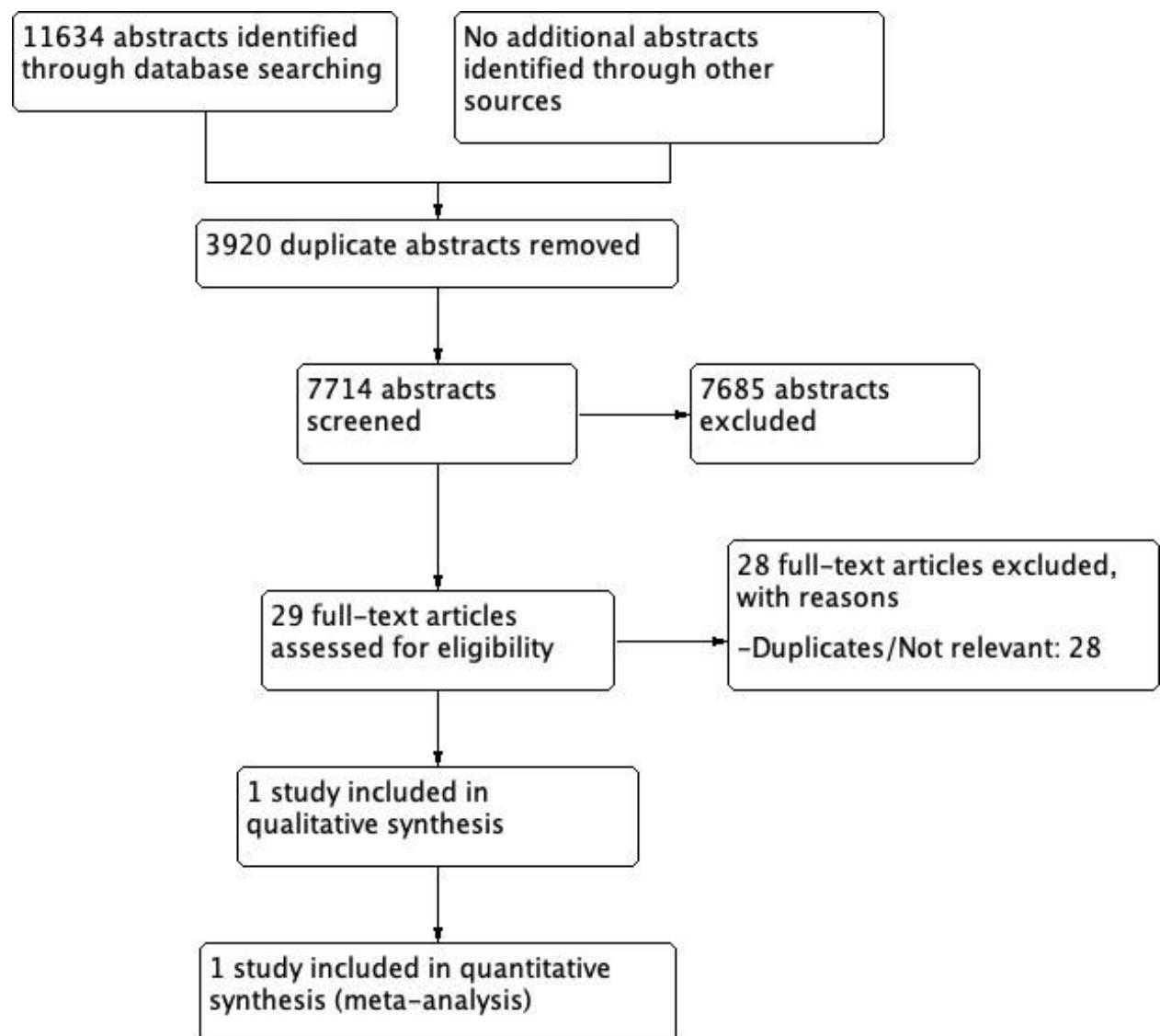
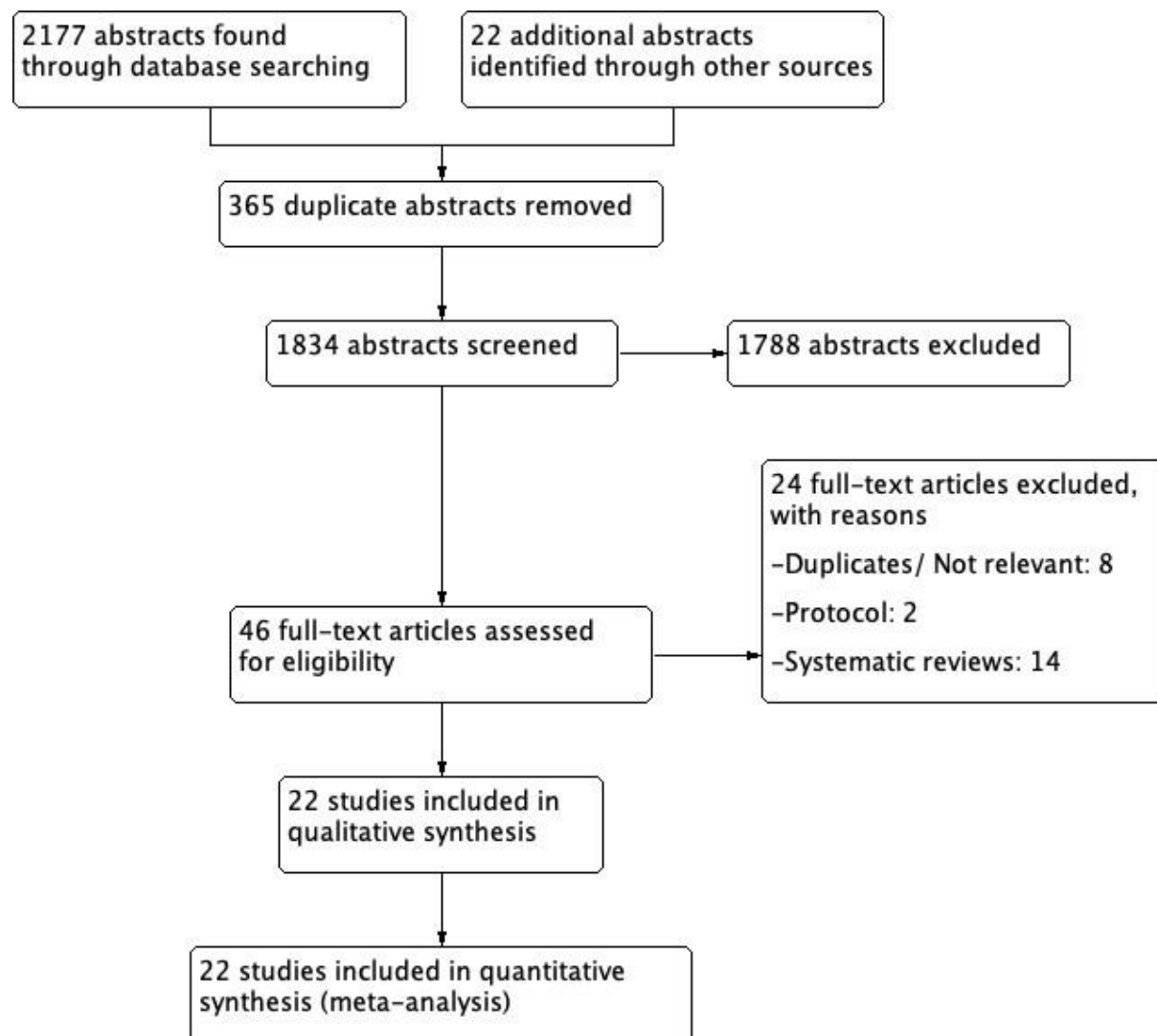


Figure S3: Literature search stenting versus endarterectomy



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