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Optimal Management of Asymptomatic Carotid Artery Stenosis: A Systematic Review and Network Meta-Analysis

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1 Short title: Systematic Review and Network Meta-Analysis of Optimal Management in
2 Asymptomatic CAS

3

4 **Optimal Management of Asymptomatic Carotid Artery Stenosis: A Systematic Review**
5 **and Network Meta-Analysis**

6

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18

19 **WHAT THIS PAPER ADDS**

20 Optimal management of asymptomatic carotid stenosis remains challenging. Best medical
21 treatment (BMT) has evolved such that it is no longer appropriate to combine different
22 medical regimens under the broad heading of BMT. Therefore, a network meta-analysis was
23 undertaken, which included traditional and modern BMT as separate arms. The results
24 suggest that modern BMT may have comparable efficacy to carotid endarterectomy in
25 reducing morbidity.

26

27 **Objective:** Management of asymptomatic carotid artery stenosis (ACAS), including carotid
28 endarterectomy (CEA), carotid artery stenting (CAS), and best medical treatment (BMT),
29 remains inconsistent in current practice. Early studies reported a benefit of CEA *vs.* BMT;
30 however, the current risk–benefit profile of invasive therapy lacks consensus. By evaluating
31 the effects of modern BMT *vs.* invasive intervention on patient outcomes, this study aimed to
32 influence the future management of ACAS.

33 **Methods:** A systematic review and series of network meta-analyses were performed
34 assessing peri-operative (within 30 days) and long term (30 days – 5 years) stroke and
35 mortality risk between ACAS interventions. Total stroke, major, minor, ipsilateral, and
36 contralateral stroke subtypes were assessed independently. Traditional (pre-2000) and
37 modern (post-2000) BMT were compared to assess clinical improvements in medical therapy
38 over the previous two decades. Risks of myocardial infarction (MI) and cranial nerve injury
39 (CNI) were also assessed.

40 **Results:** Seventeen reports of 14 310 patients with > 50% ACAS were included. CEA
41 reduced the odds of a peri-operative stroke event occurring *vs.* CAS (odds ratio [OR] 1.6,
42 95% confidence interval [CI] 1.1 – 2.2 [0 – 20 fewer/1 000]). CEA and CAS reduced the long
43 term odds of minor strokes (OR 0.35, 95% CI 0.21 – 0.59 [20 fewer/1 000]) and ipsilateral
44 strokes (OR 0.27, 95% CI 0.19 – 0.39 [30 fewer/1 000]) *vs.* all BMT. CEA reduced the odds
45 of major strokes and combined stroke and mortality *vs.* traditional BMT; however, no
46 difference was found between CEA and modern BMT. CAS reduced the odds of peri-
47 operative MI (OR 0.49, 95% CI 0.26 – 0.91) and CNI (OR 0.07, 95% CI 0.01 – 0.42) *vs.*
48 CEA.

49 **Conclusion:** Modern BMT demonstrates similar reductions in major stroke, combined
50 stroke, and mortality to CEA. The overall risk reductions are low and data were unavailable
51 to assess subgroups which may benefit from intervention. However, BMT carries the
52 potential to reduce the requirement for surgical intervention in patients with ACAS.

53

54 **Keywords:** Best medical treatment, Carotid artery disease, Carotid endarterectomy, Carotid
55 stenting, Meta-analysis

56

57 <H1>INTRODUCTION

58 The optimal interventional strategy for asymptomatic carotid artery stenosis (ACAS) remains
59 variable in current practice.¹ While a number of early randomised trials examining
60 asymptomatic disease identified significant benefits associated with carotid endarterectomy
61 (CEA) *vs.* best medical treatment (BMT),²⁻⁴ a distinct lack of consensus remains regarding
62 the potential risk to benefit profile of invasive intervention. This is largely related to
63 improvements in the delivery and constitution of BMT in recent decades,⁵ the evolution of
64 treatment paradigms following the introduction of carotid artery stenting (CAS), and the
65 relatively modest reductions in stroke risk associated with intervention.²⁻⁴

66 Indeed, since the conduct of the original trials of patients with ACAS,²⁻⁴ stroke rates,
67 for all degrees of ACAS, have reduced dramatically.⁵ This potentially diminishes the benefits
68 of asymptomatic CEA further and implies an improvement in BMT. Consequently, it has
69 been hypothesised that up to 94% of interventions for ACAS may be unnecessary.⁵
70 Additionally, randomised controlled trials RCTs; Asymptomatic Carotid Surgery Trial
71 [ACST-1], Asymptomatic Carotid Atherosclerosis Study [ACAS]) have shown that men
72 younger than 75–80 years with at least 60% ACAS have benefitted from CEA over BMT;
73 however, other subgroups, including women, have not been shown to benefit from CEA vs.
74 BMT.

75 CAS is a user dependent alternative that was introduced for patients deemed to be too
76 high risk for CEA, with rates of periprocedural stroke following CAS consistently higher
77 than those seen with CEA, despite ongoing developments in embolic protection devices.^{6,7}
78 Nevertheless, CAS is an economically attractive procedure for hospitals, being more
79 prevalent in for profit hospital settings.⁸ However, improved technologies, optimum patient
80 selection, and a plateau in operator experience carries the possibility of diminished rates of
81 periprocedural outcomes with further trials ongoing to evaluate this effect.

82 With regard to BMT, initial guidelines were ambiguous about the medical
83 management of ACAS and focused more on the degree of stenosis CEA. Medical
84 interventions, specifically antiplatelet therapy, were seen more as a peri-operative adjunct
85 rather than an independent strategy.⁹ It was not until 1998 that other important risk factors
86 such as blood pressure control, smoking cessation, lipid lowering drugs, moderate alcohol
87 consumption, and antiplatelet therapy were also strongly advocated for.¹⁰

88 The ACST-1 trial was a randomised trial of early vs. deferred endarterectomy with
89 BMT, and involved asymptomatic and recently asymptomatic patients with carotid stenosis.¹¹
90 This trial observed patients with ACAS managed with BMT alone from 1993 to 2003, and
91 captured the developing trends toward modern BMT used today. It showed that BMT at the
92 beginning of the trial (1993) included 91% antiplatelet drug use, 51% antihypertensive drug
93 use, 5% anticoagulant use, and 11% lipid lowering drug use. By 2002, use of
94 antihypertensives and lipid lowering therapies increased to 81% and 70% respectively.¹¹ By
95 2007, 88% antiplatelet drug use, 89% antihypertensive drug use, 11% anticoagulant use, and
96 80% lipid lowering drug use were reported.¹²

97 A systematic review of guidelines for the management of asymptomatic and
98 symptomatic carotid stenosis documented a lack of clarity and underuse of evidence on
99 medical treatment strategies alone to treat ACAS.¹³ Furthermore, the review highlighted

100 concerns that guidelines are based on trials of CEA *vs.* medical treatments alone in which
101 patients were randomised long before the modernisation of BMT, with no current literature
102 highlighting the efficacy of modern BMT. A large amount of high quality data has been
103 produced in the last four years, which may change the outcomes of previous meta-analyses.

104 Given this continued strategic uncertainty that exists in the asymptomatic patient
105 cohort, this network meta-analysis (NMA) aimed systematically and definitively to appraise
106 the current randomised trial literature to further inform treatment based decision making in
107 those with significant asymptomatic stenosis. This NMA allowed for the comparison of three
108 or more interventions simultaneously by combining both direct and indirect evidence across a
109 network of studies, yielding more precise estimates than a single direct or indirect evaluation.
110 Furthermore, it aimed to evaluate specifically the effects of modern BMT on patient
111 outcomes *vs.* invasive intervention.

112

113 <H1>MATERIALS AND METHODS

114 <H2>*Statement of design*

115 This analysis was prospectively registered with PROSPERO on 29 December 2021
116 (CRD42021294015) and performed in accordance with guidelines outlined by the Preferred
117 Items for Systematic Review and Meta-Analysis (PRISMA) extension statement for the
118 reporting of systematic reviews incorporating NMAs of healthcare interventions.¹⁴ Conduct
119 was based upon the PRISMA guidelines, the PICO (population, intervention, comparator,
120 outcome) framework,¹⁵ and the protocol defined in PROSPERO. Ethical approval was not
121 required as the outcome data have been published previously.

122

123 <H2>*Study eligibility*

124 Only published RCTs comparing at least two modalities of ACAS intervention (CEA, CAS,
125 and BMT alone) were included. Patients allocated to CEA and CAS also received BMT. The
126 minimum number of participants within a study was 50. All incorporated asymptomatic
127 carotid disease was evaluated using duplex ultrasonography and/or computed tomography or
128 magnetic resonance angiography (CTA and MRA, respectively). Any type of CEA including
129 traditional, eversion, or modified eversion was included. CAS with and without an embolic
130 protection device was included. BMT included antiplatelets, antihypertensives, lipid lowering
131 agents (i.e., statins), and education to modify risk factors associated with increased stroke
132 risk. Included articles were required to report at least one predefined study endpoint. English
133 language reports alone were analysed, with no time limitation set for the date of publication.

134 Studies with insufficient data to permit statistical analysis were excluded. Primary authors
135 were not contacted for missing data.

136

137 <H2>Definitions

138 *Asymptomatic lesions* were defined as image confirmed carotid stenosis $\geq 50\%$ in the absence
139 of preceding neurological symptoms, indicating a cerebrovascular event in the six months
140 prior to study enrolment.

141 *Stroke* was defined as sudden onset of acute neurologic deficits with focal signs and
142 symptoms lasting 24 hours or longer. Both ischaemic and haemorrhagic strokes were
143 included. *Major stroke* was defined as one or more of the following: (1) on the basis of
144 clinical data as at least moderate disability, with patients requiring help after the stroke; (2) a
145 modified Rankin scale (mRS) score of 3 – 5 after one month; or (3) fatal (causing death
146 directly or indirectly). These symptoms must have persisted for at least one month. *Minor*
147 *stroke* was defined as one or more of the following: (1) on the basis of clinical data as slight
148 or no residual disability, with patients able to perform most activities of daily living
149 independently; or (2) an mRS score of 0 – 2 after at least one month. These symptoms must
150 have persisted for at least one month. *Ipsilateral stroke* was defined as stroke in the vascular
151 distribution of the study carotid artery. *Transient ischaemic attack* (TIA) was defined as a
152 focal neurological deficit of abrupt onset lasting at least 30 seconds and resolving completely
153 within 24 hours.

154 *Myocardial infarction* (MI) was defined as one or more of the following occurring within
155 one month postprocedure: (1) elevated enzymes (creatine kinase, troponins); (2)
156 electrocardiographic evidence of ischaemia; or (3) chest pain or symptoms consistent with
157 ischaemia.

158 *Cranial nerve injury* (CNI) was defined as injury to the cranial nerve occurring within
159 one month postprocedure.

160 *Traditional BMT* was defined as BMT that included only antiplatelet agents. Traditional
161 BMT also included discussion of stroke risk factors and their modification, including
162 hypertension, diabetes, abnormal lipid levels, excessive alcohol consumption, and tobacco
163 use. This therapy was used in trials with recruitment dates prior to 2000. *Modern BMT* was
164 defined as BMT including combinations of antiplatelet, antihypertensive, and lipid lowering
165 agents. Modern BMT advocated for smoking cessation and moderate alcohol consumption,
166 and glycaemic control. These therapies were used in trials with recruitment dates after 2000.

167 The *peri-operative period* refers to events occurring within 30 days postprocedure (CEA
168 or CAS). *Long term* refers to events occurring between 30 days to five years.

169

170 <H2>Population, interventions, controls, and outcomes

171 Patients with ACAS established with objective diagnostic measures (i.e., duplex
172 ultrasonography, MRA, and CTA) were included in this study. Interventions assessed include
173 CEA + BMT, CAS + BMT, and BMT alone. All interventions were administered in
174 conjunction with BMT and will hereafter be referred to as CEA, CAS, or BMT. The control
175 variable was any alternative, distinct ACAS treatment modality to that described in the
176 intervention group. The primary goal was to determine which ACAS intervention
177 demonstrates the lowest risk of stroke within a 30 day peri-operative period and in the long
178 term (30 days – 5 years). Stroke subtypes included ipsilateral, contralateral, major, and minor
179 stroke. Secondary goals included determining which intervention had the lowest risk of TIA
180 and combined stroke and mortality between 30 days and five years, and the rate of adverse
181 events (AEs). AEs assessed included MI and CNI within the 30 day peri-operative period.

182

183 <H2>Search methodology and data extraction

184 MEDLINE via PubMed, Embase, Scopus, and the Cochrane Central Register of Controlled
185 Trials were systematically searched for relevant titles from 1990 until November 2021. Search
186 medical subject headings (MeSH) in English included "asymptomatic carotid artery stenosis"
187 [Mesh] AND "medical therapy" [Mesh] OR "asymptomatic carotid artery stenosis" [Mesh]
188 AND "surgery" [Mesh] OR "asymptomatic carotid artery stenosis" [Mesh] AND "stent"
189 [Mesh] for each database. Manual cross referencing of reference lists from previous reviews
190 and included studies was also undertaken.

191 Records were initially screened by abstract. Data searching and extraction were
192 completed by two authors independently, with disagreements resolved in consultation with
193 the senior author. All parts of the search and extraction were duplicated by two authors
194 independently. Extracted data included study demographic and clinicopathological
195 information, and described outcomes of interest. All data were tabulated in accordance with
196 study protocols.

197

198 <H2>Data management and analysis

199 Descriptive statistics were used to report characteristics of included trials. Rates of stroke,
200 mortality, and adverse events for each intervention were presented as odds ratios (ORs)

201 and/or risk difference (RD). ORs and RDs were calculated using crude event RCT data, to
202 compare interventions using an intention to treat analysis, where applicable. The latest
203 recorded data were analysed when multiple assessments were reported within any time frame.

204 NMAAs were conducted using the netmeta and Shiny packages for R, with a
205 frequentist statistical method, using a random effects model.¹⁶ A random effects model was
206 used to account for interstudy heterogeneity. Dichotomous meta-analyses were performed
207 using Review Manager (RevMan Version 5.4; Nordic Cochrane Centre, Copenhagen,
208 Denmark) following the Mantel–Haenszel method. Effect sizes were described with a 95%
209 confidence interval (CI). Results were considered statistically significant at a p value of $<$
210 $.050$ if the 95% CI did not include the value of 1. Rank probabilities were plotted against the
211 possible ranks for all competing treatments. Quality of the network and indirect analyses was
212 performed using p values associated with each comparison in the inconsistency tables
213 generated from the analysis.

214 To assess whether advances in BMT make it as equally efficacious as its invasive
215 counterparts, all available BMT data were analysed to identify a timepoint at which trials
216 adopted modern BMT guidelines (antiplatelet, antihypertensive, lipid lowering agents,
217 glycaemic control, and lifestyle/diet modifications) in their comparative arms. Studies after
218 2000 had all adopted modern BMT strategies and trials before 2000, unless otherwise stated,
219 used traditional BMT strategies. Therefore, BMT was subcategorised into traditional BMT
220 (pre-2000), modern BMT (post-2000), and combined BMT (all BMT data included).

221 The GRADE (Grading of Recommendations, Assessment, Development and
222 Evaluations) framework was used to assess the quality of evidence presented in this NMA.¹⁷
223 Methodological assessment of the included studies was undertaken using the Cochrane risk
224 of bias assessment tool.¹⁸

225

226 <H1>RESULTS

227 <H2>Search overview and characteristics of included studies

228 The initial systematic search identified a total of 4 646 records, which provided 2 647 records
229 for review after the removal of duplicates (Fig. 1). Forty-two full text articles were assessed,
230 with 17 records eligible for inclusion, encompassing 13 RCTs and 14 310 patients.^{2,4,6,19–30}

231 Studies were primarily conducted in North America ($n = 7$) and Europe ($n = 5$), with
232 publication dates ranging from 1990 to 2021 and reported follow up to five years. Outcome
233 data were reported in the peri-operative period ($n = 7/13$) and follow up to five years, with

234 intervals ranging from one to five years. All randomised participants underwent intervention
235 for ACAS. The pooled mean age of participants was 67.9 years (range 40 – 91 years);^{9,22} the
236 majority of patients in all reports were men. Further study characteristics are summarised in
237 Table 1 and Supplementary Table S1. Peri-operative outcomes within 30 days included CAS
238 and CEA only. BMT was added into the NMAs after the peri-operative period (beyond 30
239 days; Figs 2 and 3). Individuals allocated to the deferred CEA cohort in the ACST-1 trial
240 were classified as receiving BMT alone. The definitions for each outcome were extracted
241 directly from the included trials.

242

243 <H2>Stroke

244 <H3>Peri-operative stroke. Six RCTs (6 855 patients) provided data for peri-operative stroke
245 within a 30 day period. This was a dichotomous comparison between CEA and CAS. Five
246 studies reported data on all stroke events (Supplementary Figure S1). One study reported no
247 events in either cohort and was excluded.²⁹ CEA reduced the odds of all stroke events
248 occurring within 30 days peri-operatively vs. CAS (OR 1.6, 95% CI 1.1 – 2.2). CEA
249 specifically reduced the odds of a minor peri-operative stroke (OR 1.7, 95% CI 1.1 – 2.6)
250 (Supplementary Figure S2) but did not affect the odds of a major stroke event occurring (OR
251 0.89, 95% CI 0.33 – 2.3; Supplementary Figure S3). Neither intervention reduced the odds of
252 contralateral peri-operative stroke (Supplementary Figure S4), or ipsilateral stroke event (OR
253 1.7, 95% CI 0.96 – 3.1; Supplementary Figure S5). GRADE assessment established a
254 moderate degree of confidence to support these estimates (Table 2).

255

256 <H3>Long-term stroke. Thirteen RCTs (11 693 patients) comparing BMT alone, CAS + BMT,
257 and/or CEA + BMT reported stroke events occurring between 30 days and five years (Table
258 2). There was a moderate to high degree of certainty that CEA significantly reduced the odds
259 of all stroke events compared with BMT pre-2000 (OR 0.65, 95% CI 0.43 – 0.99) and BMT
260 post-2000 (OR 0.53, 95% CI 0.30 – 0.94; Supplementary Table S2). However, no significant
261 risk difference was established between CEA and both traditional or modern BMT (30 fewer
262 per 1 000 cases [95% CI 30 fewer – 10 more and 95% CI 30 fewer – 20 more, respectively]).

263 When compared to combined BMT, both CEA (OR 0.35, 95% CI 0.21 – 0.59) and
264 CAS (OR 0.51, 95% CI 0.26 – 0.98) significantly reduced the odds of a minor stroke
265 occurring up to five years (Supplementary Table S3). GRADE assessment established a high
266 degree of certainty that CEA resulted in 20 fewer minor stroke events per 1 000 patients
267 (95% CI –0.03 – –0.01) than modern BMT (Table 2). CAS resulted in 10 fewer minor stroke

268 events per 1 000 patients (95% CI $-0.02 - 0.0$) than modern BMT (Table 2). CEA (OR 0.27,
269 95% CI 0.19 – 0.39) and CAS (OR 0.37, 95% CI 0.22 – 0.62) also reduced the odds of an
270 ipsilateral stroke occurring up to five years *vs.* combined BMT alone, a finding consistent
271 with the use of BMT before and after 2000 (Supplementary Table S4). When compared to
272 CEA, there was a high level of certainty that traditional BMT resulted in 40 more events per
273 1 000 patients (95% CI $-0.02 - -0.06$) and modern BMT resulted in 30 more events per 1
274 000 patients (95% CI $-0.01 - -0.04$; Table 2). CEA reduced the odds of a major stroke *vs.*
275 traditional BMT (OR 0.62, 95% CI 0.38 – 1.0); however, no difference was found between
276 CEA and modern BMT (OR 0.59, 95% CI 0.32 – 1.1; Supplementary Table S5). With a low
277 degree of confidence, the results demonstrated that CEA resulted in 10 fewer major stroke
278 cases per 1 000 (95% CI $0 - -0.01$) *vs.* modern BMT (Table 2).

279

280 <H2>*Mortality*

281 <H3>**Peri-operative mortality.** Six RCTs (6 855 patients) comparing CEA and CAS reported
282 mortality occurring within the 30 day peri-operative period. Four trials were excluded from
283 the analysis due to zero events in either arm.^{6,7,26,29} Neither procedure was favoured to reduce
284 the odds of 30 day peri-operative mortality (Supplementary Figure S6).

285

286 <H3>**Long-term mortality.** Nine RCTs (11 101 patients) comparing CEA + BMT, CAS +
287 BMT, and BMT alone reported mortality from 30 days to five years. With a moderate level
288 of certainty, the results demonstrated that no treatment was significantly superior in reducing
289 the odds of mortality from 30 days to five years (Supplementary Table S6).

290

291 <H2>*Stroke and mortality*

292 Nine RCTs (4 690) reported the odds of stroke and mortality combined from 30 days to five
293 years, in patients who received CEA + BMT, CAS + BMT, or BMT alone (Fig. 3G). One
294 trial was excluded from the analysis due to zero events occurring in any treatment arm.²¹
295 GRADE assessment established a moderate degree of confidence for all results in this cohort
296 (Table 2). CEA significantly reduced the odds of stroke and mortality compared with BMT
297 therapy pre-2000 (OR 0.72, 95% CI 0.58 – 0.89); however, no significant difference was
298 found between modern BMT and CEA (OR 0.56, 95% CI 0.27 – 1.2; Supplementary Table
299 S7). CEA resulted in 40 fewer events per 1 000 patients (95% CI $-0.08 - 0.0$) *vs.* traditional
300 BMT but did not show a significant risk reduction *vs.* modern BMT (RD -0.04 , 95% CI -0.1
301 $- 0.02$; Table 2).

302

303 <H2>Trans-*ischaemic attack*

304 Four RCTs (1 513 patients), reported TIA events occurring between 30 days and five years
305 (Figure 3H). With a high degree of certainty, CEA ranked best in reducing the odds of a TIA
306 within five years, significantly reducing the odds compared with combined BMT (OR 0.58,
307 95% CI 0.35 – 0.96; Supplementary Table S8).

308

309 <H2>Adverse events

310 <H3>Myocardial infarction. Six RCTs (855 patients) assessed MI events within a 30 day peri-
311 operative period. Three RCTs were excluded from the analysis due to zero events.^{7,26,29} There
312 was moderate certainty that CAS reduced the odds of peri-operative MI vs. CEA (OR 0.49,
313 95% CI 0.26 – 0.91; Supplementary Figure S7). However, the risk difference was 0 fewer
314 cases per 1 000 (95% CI 0 – -0.01; Table 2).

315

316 <H3>Cranial nerve injury. Five RCTs (5 360 patients) reported CNI within a 30 day peri-
317 operative period. CAS reduced the odds of CNI vs. CEA (OR 0.07, 95% CI 0.01 – 0.42;
318 Supplementary Figure S8). There was moderate certainty that the risk difference was 40
319 fewer per 1 000 (95% CI 0 – -0.07; Table 2). Only one RCT reported a CNI following
320 CAS.²⁰

321 Supplementary Tables S9 – S18 provide confirmation of the statistical accuracy of the
322 analysis and can be viewed online.

323

324 <H1>DISCUSSION

325 ACAS remains a prevalent public health concern, potentially contributing to significant rates
326 of disability and death.³¹ Strokes, both major and minor, can have disabling consequences,
327 with losses of function in performing activities of daily living. CEA + BMT, CAS + BMT,
328 and BMT alone are interventions to manage ACAS that have been examined in the literature;
329 however, in the context of ever improving medical therapy, the optimal therapeutic approach
330 in this cohort remains a topic of contention. While previous trial data have continually
331 highlighted the superiority of CEA or CAS over BMT with regard to stroke prevention,^{2,4}
332 later studies have suggested that the stroke risk in asymptomatic disease may be similar
333 following the introduction of newer lipid-lowering and antiplatelet agents.^{32–34}

334 The current study showed that modern BMT and CEA + BMT showed equivalent
335 efficacy in reducing major strokes. This was a significant improvement compared with
336 cohorts treated with traditional BMT. Furthermore, no significant difference in TIA risk or
337 mortality risk was established between CEA + BMT and BMT. When comparing stroke
338 subtypes, this study found that CEA + BMT reduced ipsilateral stroke events *vs.* modern
339 BMT alone. Furthermore, CEA + BMT reduced the odds of a contralateral stroke event
340 occurring; however, this result was relative, unsupported by a negligible clinical risk
341 difference (0 – 20 fewer events per 1 000 patients).

342 The widespread use of lipid lowering drugs in the past two decades have promised
343 improved outcomes and reduced stroke risks in patients using BMT alone.^{32,35,36} In fact, in a
344 meta-regression analysis of 26 studies, Raman *et al.* found that the rate of ipsilateral stroke
345 was significantly lower in studies that closed recruitment between 2000 and 2010 (stroke risk
346 1.13% per annum) compared with those recruited before 2000 (stroke risk 2.38% per
347 annum).³⁰ To account for differences in BMT prior to 2000, each outcome was assessed as
348 combined (all studies from 1990 to 2021) traditional and modern BMT. In this NMA, when
349 combined BMT (all studies from 1990 to 2021) was analysed, CEA + BMT was favoured to
350 reduce long term stroke and most stroke subtypes and TIA *vs.* BMT. This is in agreement
351 with studies that have compared BMT to CEA.^{3,28,37} This may be attributed to a multitude of
352 factors, perhaps increased use of medical therapy. For example, in the ACST-1 trial, the use
353 of statins increased from 9% in 1993 to 81% in 2007.²⁴ Interestingly, CEA + BMT
354 significantly reduced the odds of a major stroke compared with BMT before 2000; however,
355 after 2000, this significance was lost, indicating no significant difference in treatment. This
356 trend was also seen to reduce the odds of stroke and mortality.

357 A reduction of stroke risk with modern BMT has been demonstrated in the
358 literature.^{24,35,36,38} For example, the annual ipsilateral stroke risk of BMT with aspirin alone
359 during the ACAS trial in 1995 was 2.2%,² improving to 1% in the ACST-1 trial in 2004,^{3,24}
360 when BMT included antiplatelet, antihypertensive, and lipid lowering therapies. In ACST-1,³
361 lipid lowering treatment demonstrated lower rates of long term stroke in both groups,
362 indicating that intensive medical therapy may decrease stroke risk by stabilising plaques and
363 reducing microembolisation.^{39,40} A decline in micro-emboli has coincided with better control
364 of lipids and slower progression of carotid plaques, suggesting that the lesions themselves
365 may be altered by the era of BMT.³⁹ Maintaining a healthy lifestyle and modifying risk
366 factors, including hypertension, obesity, poorly controlled diabetes, increased alcohol intake,
367 and smoking, have been associated with 90% of the risk of stroke.⁴¹ Therefore, targeted

368 interventions aimed at modifying these risk factors can substantially reduce the burden of
369 stroke.⁴¹

370 Therefore, this analysis supports the efficacy of modern BMT in reducing major
371 stroke and combined stroke and mortality. The risk reduction of stroke between CEA + BMT
372 and modern BMT are low, and current studies are reported in such a way that analysis of
373 subgroups that may benefit from intervention cannot be performed (i.e., degree of stenosis
374 and plaque morphology). Awareness that stroke risk in asymptomatic patients treated with
375 BMT may be less than previously thought has led to calls for contemporary RCTs to include
376 BMT as a separate intervention arm. Such studies should aim to document clearly the nature
377 of the current BMT utilised. BMT should consist of the diagnosis and management risk
378 factors, including hypertension, lipids, diabetes, tobacco smoking, atrial fibrillation, physical
379 inactivity, sleep disorders, and excessive weight and alcohol consumption.⁴² A blanket
380 approach to CEA may not be the optimal treatment, and risk stratification to identify high-
381 risk treatment groups may offer improvements in management as future studies continue to
382 compare these modalities.

383 CAS has been reviewed in the literature in the management of severe ACAS. In this
384 NMA, CEA reduced the odds of a stroke event *vs.* CAS within the peri-operative period, a
385 finding consistent with previous trials.^{6,7,20,25} Previous trials indicated that stroke risk was
386 primarily increased due to peri-interventional ipsilateral strokes, decreasing considerably
387 afterward.^{6,20} Despite ongoing developments in embolic protection devices, the rates of
388 periprocedural stroke following CAS remain consistently higher than those seen in CEA.^{6,7}

389 CAS + BMT was found to reduce significantly the odds of a peri-operative MI and
390 CNI compared with CEA + BMT. Although this finding is consistent with recent
391 literature,^{20,22,43} this NMA challenges the clinical relevance of these reductions in MI and
392 CNIs as they are unsupported by a negligible risk difference (0 – 10 and 0 – 70 fewer events
393 per 1 000 cases, respectively). There has been a changing environment for CAS over the last
394 two decades with advances in distal protection and interventional devices, including balloon
395 guiding catheters and more flexible catheters and wires.⁴⁴ While these ongoing advances may
396 demonstrate success in reducing procedural strokes, further comparative analyses reflective
397 of modern technology is required prior to CAS use in routine clinical practice.⁴⁵

398 Despite the systematic search strategy, some treatment comparisons in the literature
399 were limited. To date, one RCT directly compared BMT alone and CAS. Some interventions
400 were under-represented in the number of available RCTs. Underpowering of specific

401 interventions may have resulted in the absence of statistically significant differences.
402 Additionally, it is important to note that it was not possible perform sensitivity analyses
403 assessing outcomes based on plaque morphology and degree of stenosis. These analyses carry
404 the potential to identify patient cohorts more likely to benefit from intervention for ACAS
405 and should have greater emphasis in future studies. In addition, some data could not be
406 extracted due to study endpoints being combined with other outcomes. There was significant
407 heterogeneity among the reported time intervals for each outcome, resulting in results being
408 reported in wider rather than shorter time intervals. Thirdly, heterogeneity in follow up times
409 resulted in reporting wider rather than specific time frames. Fourthly, data on exact BMT
410 regimens, including resources and education, were not provided in some of the included
411 RCTs, limiting the possibility of collecting data on other aspects of BMT offering additional
412 correlations. Additionally, there was heterogeneity within the cohort treated with CAS with
413 regard to the use of technology such as embolic protection devices. Finally, this analysis was
414 limited to reports written in English and published in peer-reviewed journals; therefore, this
415 review could be affected by language and publication bias (see Table 2).

416 <H2>Conclusions

417 Modern BMT demonstrates similar reductions in major stroke, combined stroke, and
418 mortality to CEA + BMT. However, the overall risk reductions are low and current data are
419 not available to assess subgroups that may still benefit from intervention. BMT has the
420 potential to reduce the requirement for surgical intervention in patients with ACAS.

421

422 CONFLICT OF INTEREST

423 None.

424

425 FUNDING

426 None.

427

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- 555

Characteristics.

Country (centres)	Sample dates	Sample size (patients)	Interventions	BMT	Emboolic protection device, <i>n</i> (%)	Follow up duration	Degree of ICA stenosis
Austria, Germany, Switzerland (36)	2009–2014	513	CEA (203), CAS (197), BMT (113)	“As per current guidelines”	71 (36.0)	30 days–5 years	70–99%
USA and Canada (117)	2000–2008	2 502	CEA, CAS		1 213 (96.1)	4–10 years	50–69% US; ≥70% US, CTA MRA
USA and Canada (39)	1987–1993	1 662	CEA, BMT	Aspirin	–	2.7 years	≥60%
USA (65)	2005–2013	1 453	CEA (364), CAS (1 089)		1 089 (100%)	5 years	70–99% US or CT
USA (1)	1998–2002	85	CEA (42), CAS (43)		0 (0%)	4 years	>80%
USA (29)	2000–2002	237	CEA (120), CAS (117)		117 (100%)	1–3 years	≥80%
USA (11)	1983–1987	444	CEA (211), BMT (233)	Aspirin	–	3.9 years	≥50%
30 countries, 126 centres	1993–2003	3 120	CEA (1 560), BMT (1 560)	Antiplatelet, antihypertensive, lipid lowering therapies	–	5–10 years	≥60%

33 countries (130)	2008–2020	3 625	CEA (1 814), CAS (1 811)		1 344 (85% of per protocol sample)	5 years	≥60%
Israel (1)	Not reported	136	CEA (68), CAS (68)		68 (100%)	26 months	>70%
Germany (10)		410	CEA (206), BMT (204)	Aspirin and dipyridamole	–	3 years	50–90%
Russia (3)	2009–2013	55	CEA (31), BMT (24)	Antiplatelet (ASA), lipid lowering (atorvastatin), antihypertensive (amlodipine/losartan/hydrochlorothiazide) therapy	–	5 years	70–79%
USA (1)	2011–2013	60	CEA (31), CAS (29)		29 (100%)	6 months	>80%

RCT = randomised controlled trial; BMT, best medical treatment; ICA = internal carotid artery; SPACE-2 = Stent-Protected Angioplasty versus Carotid Endarterectomy at 2 years; CEA = carotid endarterectomy; CAS = carotid artery stenting; US = ultrasound; CREST = Carotid Revascularisation Endarterectomy Versus Stenting Trial; CTA = computed tomography angiography; MRA = magnetic resonance angiography; ACAS = Asymptomatic Carotid Atherosclerosis Study; ACT-1 = Asymptomatic Carotid Trial; ACST-1 = Asymptomatic Carotid Surgery Trial; ACST-2 = Asymptomatic Carotid Surgery Trial 2; AMTEC = aggressive medical treatment evaluation for asymptomatic carotid artery stenosis; ASA = acetylsalicylic acid.

Table 2. GRADE (Grading of Recommendations, Assessment, Development and Evaluations) analysis rank.		
Outcome	Effects and confidence in the estimate of effects	
	CAS	
Long Term Stroke: Total (Up to 5 Years)		No. Trials: 6
<i>Follow Up: Range from</i>	<i>26 - 120 Months</i>	No. Patients: 8
CEA	OR 0.76 [0.51; 1.15] Network estimate	10 fewer per 1 <i>more)*</i>
	Confidence of estimate: ⊕⊕⊕○	Moderate due to
Rank 1/5	Rank 2/5	
Long Term Stroke: Major (Up to 5 Years)		No. Trials: 3
<i>Follow Up: Range from</i>	<i>32 - 120 Months</i>	No. Patients: 4
CEA	OR 0.91 [0.54; 1.55] Network estimate	0 fewer per 1 <i>more)*</i>
	Confidence of estimate: ⊕⊕○○	Low due to impr
Rank 1/5	Rank 2/5	
Long Term Stroke: Ipsilateral (Up to 5 Years)		No. Trials: 6
<i>Follow Up: Range from</i>	<i>6 - 120 Months</i>	No. Patients: 6
CEA	OR 0.75 [0.50; 1.12] Network estimate	10 fewer per 1 <i>fewer)*</i>
	Confidence of estimate: ⊕⊕⊕○	Moderate due in
Rank 1/5	Rank 2/5	
Stroke & Mortality		No. Trials: 4
<i>Follow Up: Range from</i>	<i>26 - 120 Months</i>	No. Patients: 1

CEA	OR 0.78 [0.57; 1.06] Network estimate	20 fewer per 1000 (more)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 1/5	Rank 2/5	
Mortality (30 days to 5 years)		No. Trials: 5
<i>Follow Up: Range from</i>	<i>26 - 120 Months</i>	No. Patients: 5000
CEA	OR 0.93 [0.79; 1.08] Network estimate	0 fewer per 1000 (more)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 1/5	Rank 5/5	
Long Term Stroke: Minor (Up to 5 Years)		No. Trials: 3
<i>Follow Up: Range from</i>	<i>48 - 120 Months</i>	No. Patients: 4000
CEA	OR 0.70 [0.45; 1.07] Network estimate	10 fewer per 1000 (fewer)*
	Confidence of estimate: ⊕⊕○○	Low due to Individual Study
Rank 1/3	Rank 2/3	
Long Term Stroke: Contralateral (Up to 5 years)		No. Trials: 4
<i>Follow Up: Range from</i>	<i>48 - 120 Months</i>	No. Patients: 5000
CEA	OR 0.79 [0.48; 1.29] Network estimate	10 fewer per 1000 (more)*
	Confidence of estimate: ⊕⊕○○	Low due to imprecision
Rank 1/3	Rank 2/3	
TIA (30 days to 5 years)		No. Trials: 2
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 5000

CEA	OR 0.91 [0.31; 2.65] Network estimate	10 fewer per 1000 (more)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 1/5	Rank 2/5	
Minor Stroke (Within 30 days)		No. Trials: 3
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 500
CEA	OR 0.58 [0.38; 0.88] Network estimate	10 fewer per 1000 (fewer)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 1/2	Rank 2/2	
Peri-Operative Stroke (Within 30 Days)		No. Trials: 5
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 600
CEA	OR 0.64 [0.46; 0.88] Network estimate	10 fewer per 1000 (fewer)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 1/2	Rank 2/2	
Myocardial Infarction		
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 600
CAS	OR 0.49 [0.26; 0.91] Network estimate	0 fewer per 1000 (fewer)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 2/2	Rank 1/2	
Cranial Nerve Injury		No. Trials: 5
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 500
CAS	OR 0.07 [0.01; 0.42] Network estimate	40 fewer per 1000 (fewer)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 2/2	Rank 1/2	
Mortality (Within 30 days)		No. Trials: 6
<i>Follow Up: Range from</i>	<i>0 - 30 Days</i>	No. Patients: 600

CEA	OR 0.75 [0.32; 1.75] Network estimate	0 fewer per 1000 (95% CI: 0 fewer to 100 fewer)*
	Confidence of estimate: ⊕⊕⊕○	Moderate due to imprecision
Rank 2/2	Rank 1/2	

Red boxes indicate statistically insignificant findings; green boxes indicate statistically significant findings.

FIGURE LEGEND

Figure 1. Flowchart of the literature search according to the Preferred Items for Systematic Review and Meta-Analysis (PRISMA) statement. RCT = randomised controlled trial.

Figure 2. Forest plots comparing carotid endarterectomy (CEA) to all other interventions using combined direct and indirect evidence for all outcomes. Results are presented using odds ratios (ORs) and 95% confidence intervals (CIs) with a random effects model. Outcomes included (A) all stroke events, (B) minor stroke, (C) major stroke, (D) contralateral stroke, (E) ipsilateral stroke, (F) mortality, (G) stroke and mortality, and (H) transient ischaemic attack. BMT = best medical treatment; CAS = carotid artery stenting.

Figure 3. Network plots of all possible pairwise comparisons assessing outcomes. Outcomes included (A) all stroke events, (B) minor stroke, (C) major stroke, (D) contralateral stroke, (E) ipsilateral stroke, (F) mortality, (G) stroke and mortality, and (H) transient ischaemic attack. The size of the nodes and the thickness of edges depended on the number of people randomised and the number of trials conducted, respectively. BMT = best medical therapy; CAS = carotid artery stenting; CEA = carotid endarterectomy.

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Figure 1.

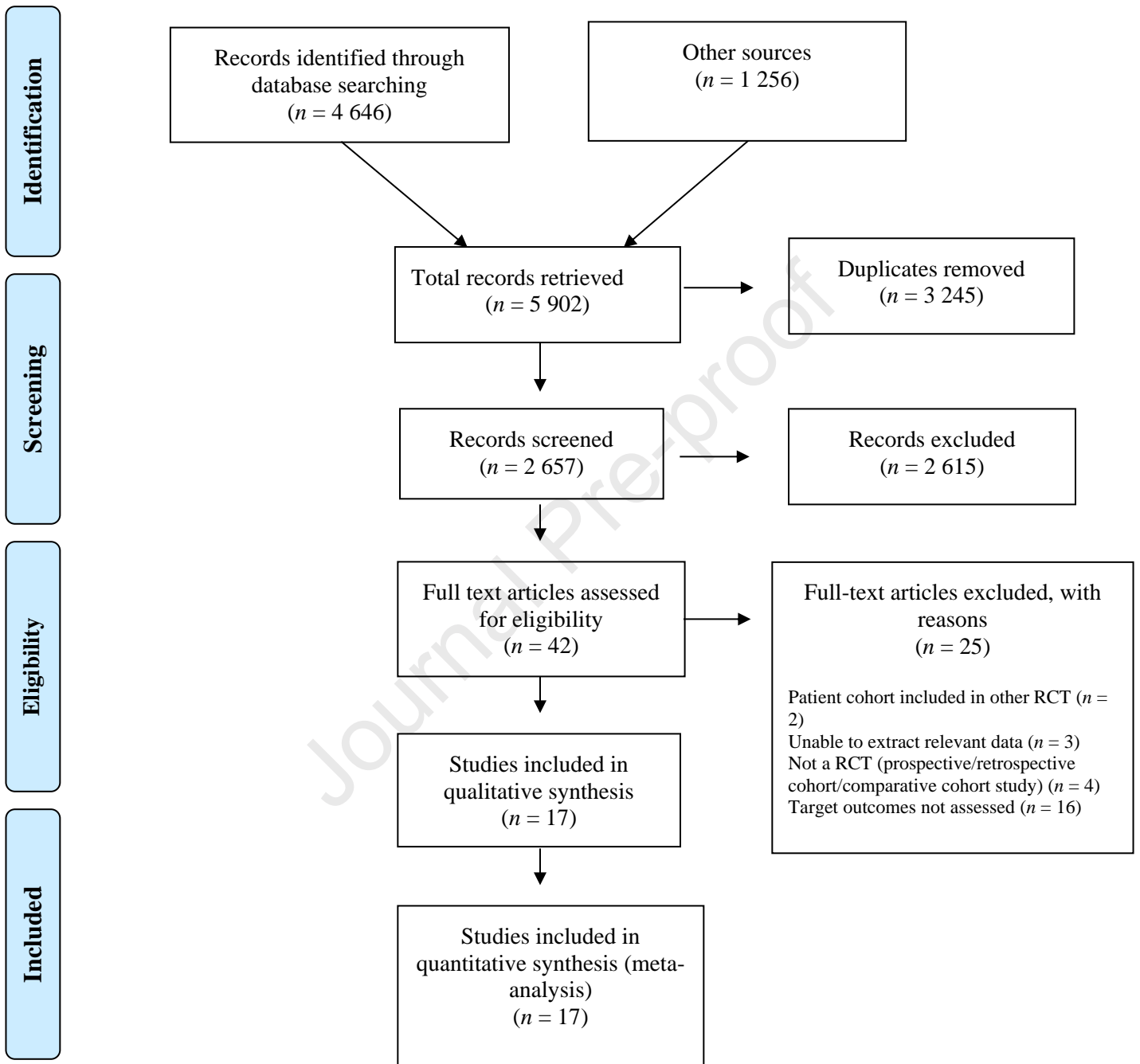


Figure 2.

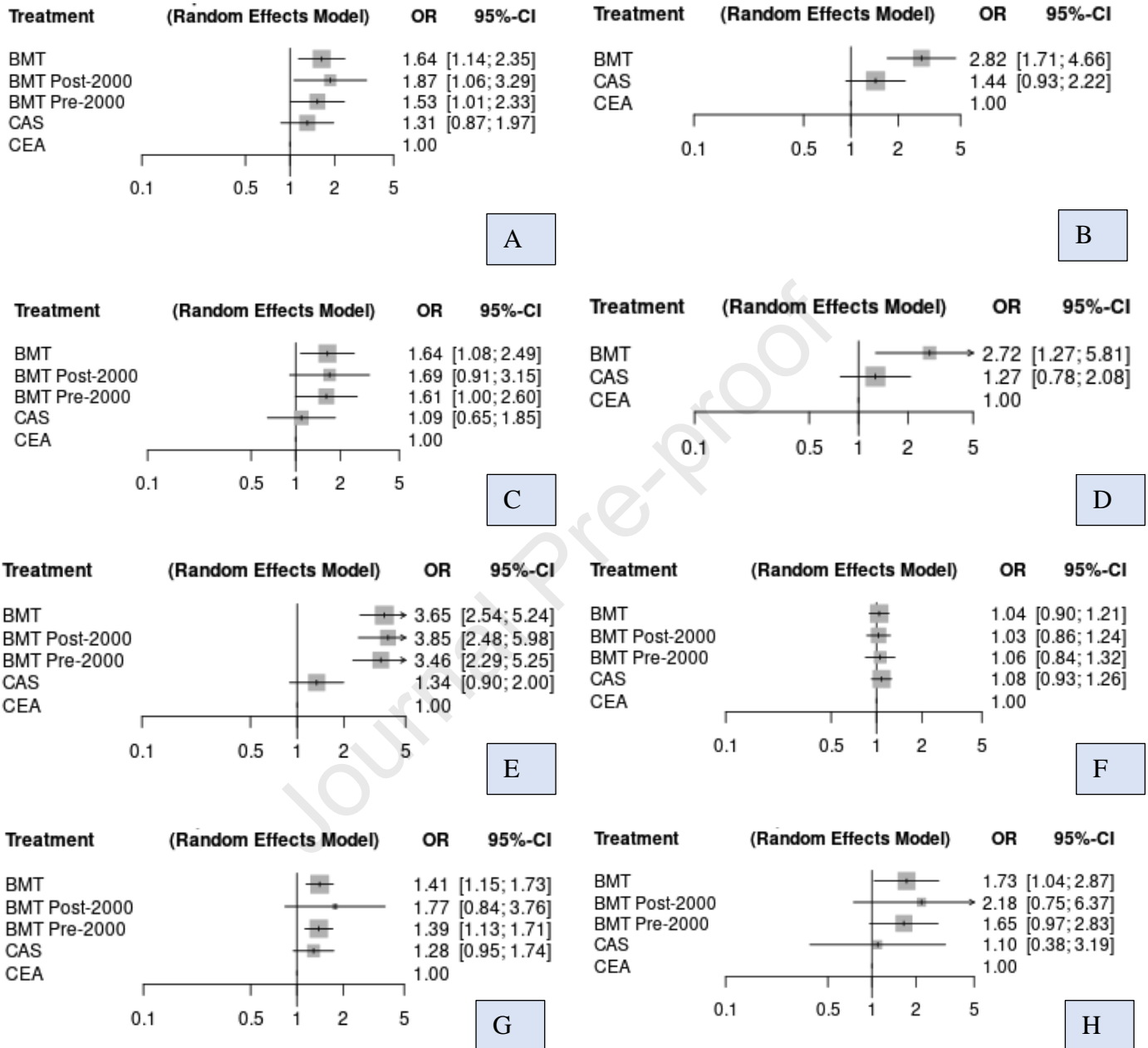
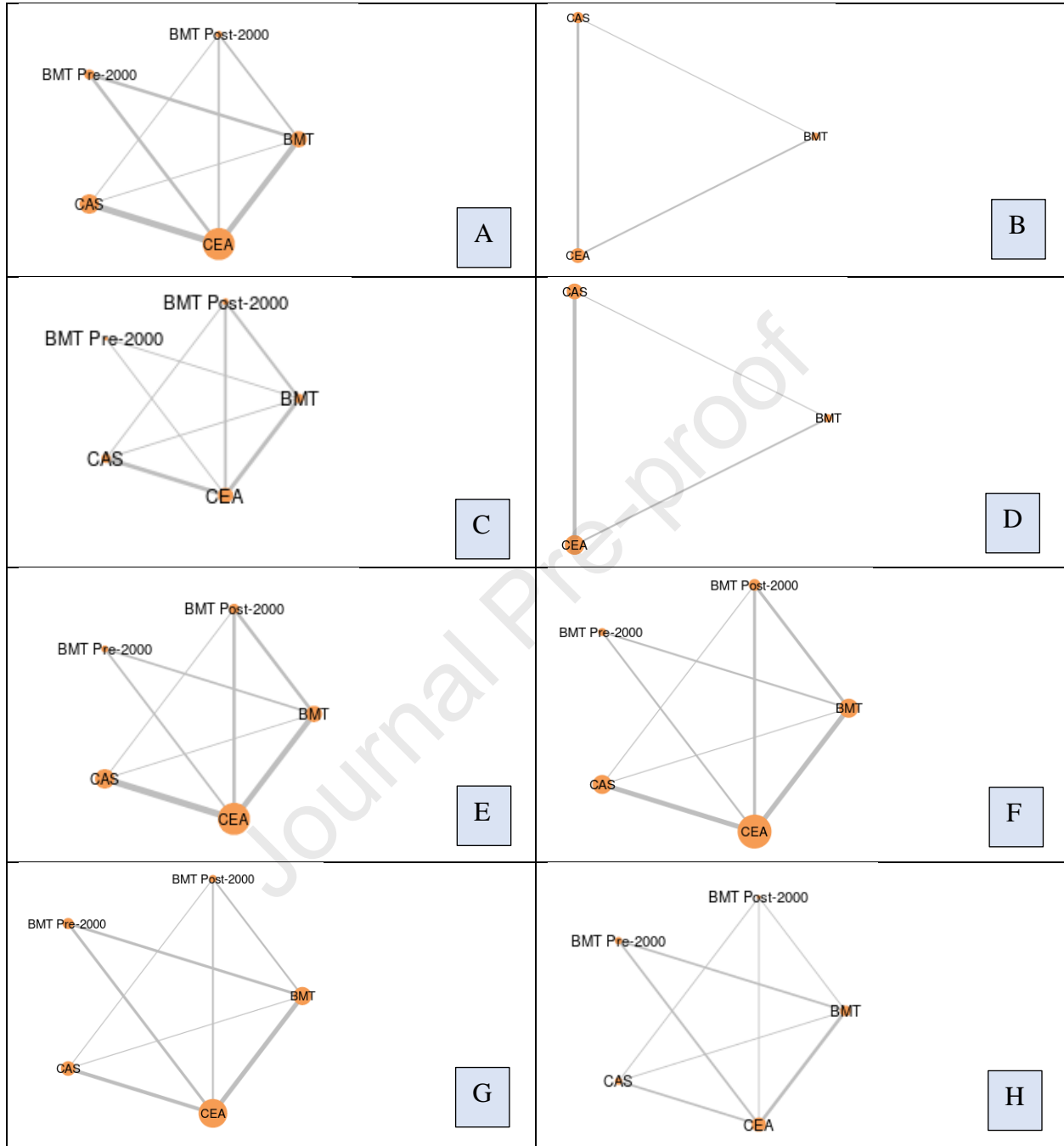


Figure 3. Network Plots of Each Comparison.



Short title

Systematic Review and Network Meta-Analysis of Optimal Management in Asymptomatic CAS

Supplementary Material

Supplementary Tables S1 – S18

Supplementary Figures S1 – S8

Figures

Figure 1 – please follow instructions on page D of guide for a PRISMA flowchart.

Figure 2 – Forest plots, please follow the instructions on page G1 onwards.

Figure 3 – other type of plot.