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Sex differences in outcome after carotid revascularization in symptomatic and asymptomatic carotid artery stenosis

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ABSTRACT

Objective: Sex differences regarding the safety and efficacy of carotid revascularization in carotid artery stenosis have been addressed in several studies with conflicting results. Moreover, women are underrepresented in clinical trials, leading to limited conclusions regarding the safety and efficacy of acute stroke treatments.

Methods: A systematic review and meta-analysis was performed by literature search including four databases from January 1985 to December 2021. Sex differences in the efficacy and safety of revascularization procedures, including carotid endarterectomy (CEA) and carotid artery stenting (CAS), for symptomatic and asymptomatic carotid artery stenoses were analyzed.

Results: Regarding CEA in symptomatic carotid artery stenosis, the stroke risk in men (3.6%) and women (3.9%) based on 99,495 patients (30 studies) did not differ (P = .16). There was also no difference in the stroke risk by different time frames up to 10 years. Compared with men, women treated with CEA had a significantly higher stroke or death rate at 4 months (2 studies, 2565 patients; 7.2% vs 5.0%; odds ratio [OR], 1.49; 95% confidence interval [CI], 1.04-2.12; $I^2 = 0\%$; P = .03), and a significantly higher rate of restenosis (1 study, 615; 17.2% vs 6.7%; OR, 2.81; 95% CI, 1.66-4.75; P = .0001). For CAS in symptomatic artery stenosis, data showed a non-significant tendency toward higher peri-procedural stroke in women, whereas for asymptomatic carotid artery stenosis, data based on 332,344 patients showed that women (compared with men) after CEA had similar rates of stroke, stroke or death, and the composite outcome stroke/death/myocardial

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infarction. The rate of restenosis at 1 year was significantly higher in women compared with men (1 study, 372 patients; 10.8% vs 3.2%; OR, 3.71; 95% Cl, 1.49-9.2; P = .005). Furthermore, CAS in asymptomatic patients was associated with low risk of a postprocedural stroke in both sexes, but a significantly higher risk of in-hospital myocardial infarction in women than men (8445 patients, 1.2% vs 0.6%; OR, 2.01; 95% Cl, 1.23-3.28; $\hat{P} = .005$).

Conclusions: A few sex-differences in short-term outcomes after carotid revascularization for symptomatic and asymptomatic carotid artery stenosis were found, although there were no significant differences in the overall stroke. This indicates a need for larger multicenter prospective studies to evaluate these sex-specific differences. More women, including those aged over 80 years, need to be enrolled in randomized controlled trials, to better understand if sex differences exist and to tailor carotid revascularization accordingly. (J Vasc Surg 2023;78:817-27.)

Keywords: Carotid endarterectomy; Carotid stenting; Ischemic stroke; Outcome; Sex differences

Stroke and transient ischemic attack (TIA) due to atherosclerotic carotid artery disease accounts for around 15% of all cases according to the definition of stroke etiology and stenosis classification used.¹ In the Caucasian population, the prevalence of carotid atherosclerotic disease, defined as \geq 50% stenosis of the carotid arteries, increases with age and is higher for men.²

Women have a higher risk of stroke during and after menopause, probably due to changes in the vascular microstructure with increasing arterial stiffness and a higher risk of hypertension.³ Recurrent carotid artery stenosis after revascularization is more prevalent in women.⁴ Carotid plaque morphology is different in women compared with men, who show higher percentages of intraplaque hemorrhage and larger necrotic cores.⁵

Sex differences in anatomy with a smaller diameter of the carotid artery in women and sex-specific risk factors during interventions can affect outcome.⁶ Biological differences, including hormonal changes, are not well-studied and likely contribute to sex differences in outcome after carotid revascularization.⁷

Moreover, women are underrepresented in randomized controlled trials (RCTs), leading to conflicting results and low evidence for interventions in women.⁸

Therefore, the aim of this study was to investigate sex differences in the efficacy and safety of revascularization procedures, including carotid endarterectomy (CEA) and carotid artery stenting (CAS), for symptomatic and asymptomatic carotid artery stenoses by performing a systematic review and meta-analysis.

METHODS

A professional methodologist (AL) prepared and executed search algorithms and strategies in four databases (MEDLINE, EMBASE, CINAHL, SCOPUS) using a combination of controlled vocabulary, free-text terms, and their corresponding Medical Subject Heading terms (Supplementary Appendix 1, online only). Potentially eligible RCTs, meta-analyses, and observational studies were identified, and citations were loaded on COVI-DENCE software. Only original articles in English from January 1985 to December 2021 were included.

The selection of studies was performed by two members of the group independently, according to predefined inclusion/exclusion criteria (Supplementary Appendix 1, online only). In case of conflict, the disagreement was resolved by a third member.

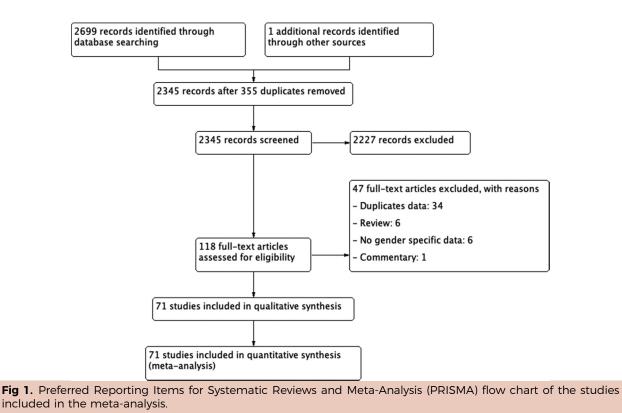
The relevant outcomes for both CEA and CAS of symptomatic and asymptomatic carotid artery stenoses selected were: ischemic stroke, TIA, mortality, myocardial infarction (MI) and/or cardiac heart failure, cranial nerve palsy, and complications of revascularization: reintervention and restenosis. After screening the titles and abstracts, the full text of potentially relevant studies was loaded onto the software and assessed following the same inclusion/exclusion criteria. The selection process is shown in the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) chart (Fig 1; for checklist, see Supplementary Appendix 2, online only). Sex-specific relevant data were extracted from eligible studies, and patients' outcomes were compared between the sexes. Due to the lack of sex-specific data in most RCTs, observational studies were also included.

Where applicable, meta-analyses were performed by using the RevMan software, using a random-effects model. Odds ratio (OR) was calculated for dichotomous variables and mean differences for continuous variables, along with their 95% confidence intervals (CIs). A value of P < .05 was considered for statistical significance. The heterogeneity was checked by a high value of I^2 and P < .05.

RESULTS

Symptomatic carotid artery stenosis. Our metaanalysis, based on 99,495 patients (35,160 women, 64,335 men) with symptomatic CAS (5 RCTs [NASCET, ECST, CREST, SPACE, CAVATAS] and 25 observational studies) treated with CEA demonstrated that the overall stroke risk did not differ between men (3.6%) and women (3.9%) (OR, 1.07; 95% CI, 0.97-1.17; $I^2 = 14\%$; P = .16) (Fig 2, A). There was also no difference in the stroke risk by different timeframes (Fig 2, A; Supplementary Table I [online only]).⁹⁻³²

The overall death rate based on 87,163 patients (31,021 women, 56,142 men) was not significantly different between women (1.5%) and men (1.4%) (OR, 0.95; 95% Cl, 0.80-1.12; $l^2 = 28\%$; P = .53) (Supplementary Fig 1, online only), whereas the death rate at 10 years was greater in men (27.1% vs 37.8% in men; P = .006) (Supplementary Table I, online only and



Supplementary Fig 1, online only).³³ Compared with men, women treated with CEA had a significantly higher stroke or death rate at 4 months (2 studies, 2565 patients; 7.2% vs 5.0%; OR, 1.49; 95% CI, 1.04-2.12; $l^2 = 0\%$; P = .03), and a slightly longer mean hospital stay (2 studies, 21,117 patients; 6.4 days vs 5.8 days; OR, 0.52; 95% CI, 0.21-0.83; $l^2 = 0\%$; P = .001).^{14,34} Women had a significantly higher rate of restenosis compared with men at both 5 years (1 study, 615 patients; 11.4% vs 3.3% in men; OR, 3.79; 95% CI, 1.89-7.61; P = .0002) and 10 years (17.2% vs 6.7%; OR, 2.81; 95% CI, 1.66-4.75; P = .0001).³³ A higher rate of cranial nerve palsy as post-procedural complication was found in women (1 study, 821 patients; 8.2% vs 4.3%; OR, 1.98; 95% CI, 1.08-3.64; P = .03) (Supplementary Table I, online only).

Regarding CAS of symptomatic carotid artery stenosis, the Carotid Revascularization Endarterectomy vs Stenting Trial (CREST) did not find a significant sex-related difference by treatment in primary endpoint rates at 4 years (P = .34).³⁵ The Stent-Protected Angioplasty vs Carotid Endarterectomy (SPACE) trial showed a non-significant increase in the periprocedural ipsilateral stroke/death for women with symptomatic carotid artery stenosis (who accounted for 28% of enrolled patients) after CAS in the subgroup analyses stratified by sex: 8.2% vs 6.4% in men (P = .48), in the CAS arm and 6.6% vs 6.0% in men (P = .85) in the CEA arm.³⁶ Based on the results of our meta-analyses, the overall stroke rate of 4650

patients (1703 women, 2947 men) did not differ between men (7.6%) and women (8.0%) receiving CAS (OR, 1.04; 95% CI, 0.79-1.38; $l^2 = 18\%$; P = .77) (Fig 2, *B*). This trend was consistent for the in-hospital stroke rate (P = .67) and for the stroke rate at 1 month (P = .28), 2 years (P = .58), and 4 years (P = .08) from stenting^{10,25} (Supplementary Table II, online only).

The risk of death (n = 7405; 2477 women, 4928 men) was also comparable between men and women (OR, 1.04; 95% CI, 0.66-1.64; l^2 = 31%; *P* = .87) (Supplementary Fig 2, online only), as well as stroke or death (n = 9615) (OR, 1.09; 95% CI, 0.91-1.30; l^2 = 0%; *P* = .37).^{17-19,23,29,36-40}

Asymptomatic carotid artery stenosis. Regarding CEA for asymptomatic carotid artery stenosis, we included in our analysis sex-specific data from 5 RCTs (ACAS, ACST, ACST 2010, CREST, ACST 2) and 17 observational studies.^{9,11,12,15,17,18,20-24,35,41-50} Overall, compared with men, women had similar rates of stroke (21 studies, 332,344 patients [144,022 women, 188,322 men]; 0.9% vs 0.8%; OR, 1.12; 95% CI, 0.96-1.30; $l^2 = 42\%$; P = .14) (Fig 3, A) and of the composite endpoint stroke/death/MI (3 studies, 5675 patients; 3.4% vs 3.2%; OR, 1.17; 95% CI, 0.75-1,3; $l^2 = 33$; P = .49).

Although the overall risk of death was slightly significantly lower in women than in men (13 studies, 313,453 patients [136,760 women, 176,693 men]; 0.35% vs 0.42%; OR, 0.87; 95% CI, 0.78-0.98; $l^2 = 0\%$; P = .02), the overall

	Wom	en	Me	n		Odds Ratio		Odds Ratio	
tudy or Subgroup	Events		Events		Weiaht	M-H, Random, 95% CI	M	-H, Random, 95% CI	
.1.1 Hospital, Symptomatic							M		
sdas 2012	66	2184	79	2184	6.4%	1.12 [0.82, 1.52]		-	
enyhel 2011, > 75 Y	121	6078	-	11408	9.7%	1.20 [0.95, 1.51]		-	
lenyhel 2011 < 75 Y	142	8887	381		11.8%	0.85 [0.70, 1.03]			
ockman 2011, Symptomatic	35	1037	30	1449	3.0%	1.65 [1.01, 2.71]			
arac 2002, Sympotmatic	12	361	20	688	1.5%	1.15 [0.55, 2.38]			
ubtotal (95% CI)		18547		36038	32.3%	1.11 [0.89, 1.38]		•	
otal events	398		700					ſ	
eterogeneity: Tau ² = 0.03; Cl		. df = 4		5): f ² = 5	7%				
est for overall effect: Z = 0.91									
		-,							
.1.2 1 month, Symptomatic									
amowitch 2005	40	753	93	1810	4.6%	1.04 [0.71, 1.52]		<u> </u>	
onati 2010	5	31	13	76	0.6%	0.93 [0.30, 2.88]			
REST 2011, Howard	6	226	15	427	0.9%	0.75 [0.29, 1.96]			
awley 2000	ě	312	9	688	0.7%	1.48 [0.52, 4.19]			
uzman 2014	4	243	9	438	0.6%	0.80 [0.24, 2.62]			
m 2014	43	960	56	1494	4.2%				
apral 2000	34	506	64	11494	3.8%				
• • • • • • •						1.21 [0.79, 1.86]		τ-	
uy 2014, Symptomatic	490	8254		12112	17.6%	1.08 [0.96, 1.22]		T I	
ine 2003, Symptomatic	4	57	3	104	0.3%	2.54 [0.55, 11.77]			
e 2003, Symptomatic	6	315	13	496	0.8%	0.72 [0.27, 1.92]			
bke 2015, Symptomatic	4	128	11	328	0.6%	0.93 [0.29, 2.97]			
attos 2001, Symptomatic	1	311	4	492	0.2%	0.39 [0.04, 3.54]			
axwell 1990, < 75 Y	7	265	2	366	0.3%	4.94 [1.02, 23.96]			
axwell 1990, > 75 Y	1	70	2	63	0.1%	0.44 [0.04, 5.00]			
lles 1994	27	838	39	1527	2.9%	1.27 [0.77, 2.09]		+	
ockman 2001, Symptomatic	20	659	25	1041	2.1%	1.27 [0.70, 2.31]		+	
chneider 1997	5	155	4	271	0.5%	2.23 [0.59, 8.41]			
slvgoulis 2014, Symptoma	4	96	4	69	0.4%	0.71 [0.17, 2.93]			
else 2004	5	56	ß	156	0.6%	1.81 [0.57, 5.80]			
ubtotal (95% CI)	-	14235	~	23098	42.0%	1.10 [1.00, 1.22]			
otal events	712		1043			• • • •		ſ	
eterogeneity: Tau ² = 0.00; Cl	-	6 df = 1		1 881- P	- 05				
est for overall effect: Z = 1.97					•				
.1.3 1 year, Symptomatic									
ark 2008	0	40	1	E 2	0.14	0 43 10 02 10 001			
ubtotal (95% CI)	0	40	1	53 53	0.1× 0.1%	0.43 [0.02, 10.89]			
		40		55	0.1%	0.43 [0.02, 10.89]			
otal events	0		1						
eterogeneity: Not applicable	/m								
est for overall effect: Z = 0.51	(P = 0.6)	1)							
1.4.2									
.1.4 2 years, Symptomatic									
ckstein 2008, SPACE	11	167	39	422	1.6%	0.69 [0.35, 1.39]		-+	
chnekder 1997	6	155	6	271	0.6%	1.78 [0.56, 5.61]			
ubtotal (95% CI)	A second s	322		693	2.2%	0.99 [0.40, 2.42]		-	
otal events	17		45						
eterogeneity: Tau ² = 0.21; Cl	n ² = 1.90	df = 1	(P = 0.1)	7); i ² = 4	7%				
est for overall effect: Z = 0.02									
154 years Symptomatic									
1.5 4 years, Symptomatic			~~						
REST 2011, Howard	14	226	23	427	1.6%	1.16 [0.58, 2.30]		<u> </u>	
ong 2016	15	102	12	171	1.2%	2.28 [1.02, 5.10]			
ubtotal (95% CI)		328		598	2.8%	1.57 [0.81, 3.05]		-	
otal events	29		35	-					
eterogeneity: Tau ² = 0.08; Cl			(P = 0.2)	1); F = 3	7%				
est for overall effect: Z = 1.35	(P = 0.1)	8)							
1.6 5 years, Symptomatic									
amowitch 2005	64	424	143	1012	6.1%	1.08 [0.79, 1.49]		+	
ine 2003, Symptomatic	4	57	3	104	0.3%	2.54 [0.55, 11.77]		<u> </u>	
ubtotal (95% CI)		481	-	1116	6.5%	1.18 [0.71, 1.94]		-	
otal events	68		146					-	
eterogeneity: Tau ² = 0.05; Cl		df = 1		8); f ² = 1	13%				
est for overall effect: Z = 0.63									
1.7 10 years, Symptomatic									
unningham 2002	22	478	81	1250	3.1%	0.70 [0.43, 1.13]		+	
attos 2001, Symptomatic	40	311	94	492	4.3%	0.62 [0.42, 0.93]			
ASCET 1999, Ferguson	76	418	166	997	6.8%	1.11 [0.82, 1.50]		-	
ubtotal (95% CI)		1207		2739	14.1%	0.81 [0.55, 1.19]		•	
otal events	138		341						
eterogeneity: Tau ² = 0.08; Cl		. df = 2		5): I ² = 6	7%				
est for overall effect: Z = 1.08	(P = 0.2	8)	0.0						
	→ v.z	~							
otal (95% CI)		35160		64335	100.0%	1.07 [0.97, 1.17]		k	
	1362		2211						
otal events	1362	a df - 4	2311	241.12	- 144				
	nt ² = 38.4		-).24); i ² (- 14%		0.02 0.1	1 10 Women Men	

Fig 2. Stroke in men and women after carotid endarterectomy (CEA) **(A)** and stenting (CAS) **(B)**. *CI*, Confidence interval; *DWI*, diffusion-weighted imaging; *MRI*, magnetic resonance imaging; *OR*, odds ratio; *Y*, years.

В		Wom	en	Mer	n		Odds Ratio	Odds Ratio
	Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI
-	2.1.1 Hospital, Symptomatic							
	Bisdas 2012	19	233	13	233	11.7%	1.50 [0.72, 3.12]	
	Dolg 2016 (1)	3	34	15	81	4.2%	0.43 [0.11, 1.58]	
	Rockman 2011, Symptomatic Subtotal (95% CI)	10	1 62 429	6	1 96 510	7.5X 23.4%	1.55 [0.60, 4.01] 1.16 [0.59, 2.30]	
	Total events	32		36				
	Heterogeneity: $Tau^2 = 0.13$; Ch Test for overall effect: $Z = 0.43$			2 (P = 0.	.22); ř	- 34%		
	2.1.2 1 month, Symptomatic							
	Bonati 2010	20	37	42	87	10.7%	1.26 [0.58, 2.73]	
	CREST 2009, Howard	7	142	14	272	7.8%	0.96 [0.38, 2.42]	
	Dua 2016 (2)	0	0	0	0		Not estimable	
	Jim 2014, Symptomatic	30	587	71	945	23.6%	0.66 [0.43, 1.03]	
	Salinas-Aragón 2016, Symp	1	30	2	81	1.3%	1.36 [0.12, 15.59]	
	Shobha 2012	2	67	4	166	2.5%	1.42 [0.25, 7.91]	
	Subtotal (95% CI)		863		1573	45.9%	0.83 [0.59, 1.16]	•
	Total events	60		133				
	Heterogeneity: $Tau^2 = 0.00$; Ch Test for overall effect: $Z = 1.09$			4 (P = 0.	.60); I ²	- 0%		
	2.1.3 2 years, Symptomatic							
	Eckstein 2008, SPACE Subtotal (95% CI)	14	171 171	42	436 436	14.6% 14.6%	0.84 [0.44, 1.57] 0.84 [0.44, 1.57]	
	Total events Heterogeneity: Not applicable Test for overall effect: $Z = 0.55$	14 (P = 0.5	6)	42				
	2.1.4 4 years, Symptomatic							
	CREST 2011, Howard Subtotal (95% CI)	23	240 240	25	428 428	16.1X 16.1%	1.71 [0.95, 3.08] 1.71 [0.95, 3.08]	
	Total events Heterogeneity: Not applicable Test for overall effect: $Z = 1.78$	23 i (P = 0.0	6)	25				
	Total (95% CI)		1703		2947	100.0%	1.04 [0.79, 1.38]	•
	Total events	129		236				
	Heterogeneity: Tau ² = 0.04; Ch Test for overall effect: Z = 0.29 Test for subgroup differences: (<u>Footnotes</u> (1) Symptomatic, DWI-MRI lesio (2) Symptomatic: OR: 12.59; 95	l (P = 0.7 Chl ² = 4.1 ns: IRR: 0	7) 83, df .49; 9	• 9 (P = 0 = 3 (P = 5% CI 0.3	0.18),	r² = 37.9	X	0.1 0.2 0.5 1 2 5 10 Women Men
Fi	g 2. Continued.							

risk of the composite endpoint stroke or death resulted slightly higher in women (8 studies, 65,340 patients; 2.0% vs 1.8%; OR, 1.30; 95% CI, 1.05-1.63; $l^2 = 39\%$; P = .02) (Fig 3, *B*).

Similarly, the rates of in-hospital MI (1 study, 49,042 patients; OR, 1.48; 95% CI, 1.17-1.85; P = .0008) and of the composite outcome stroke/MI/death (1 study, 463 patients; 5.3% vs 1.6%; OR, 3.43; 95% CI, 1.10-10.69; P = .03) were significantly higher in women than in men (Supplementary Table III, online only). However, data on these outcome measures should be interpreted with caution because they come from one study each. Perioperative (1-month) outcome events in terms of stroke (12 studies, 218,116 patients; 0.7% vs 0.6%; OR, 1.19; 95% CI, 1.01-1.40; $I^2 = 9\%$; P = .03), stroke or death (5 studies, 10,218 patients; 3.2% vs 2.1%; OR, 1.44; 95% CI, 1.13-1.85; $I^2 = 0\%$; P = .004) occurred more frequently in women than in men, except for the composite

outcome stroke/death/MI (2 studies, 4625 patients; 3.1% vs 3.2%; OR, 0.96; 95% CI, 0.69-1.34, $I^2 = 0\%$; P = .81) (Supplementary Table III, online only). The rate of restenosis at 1 year was significantly higher in women compared with men (1 study, 372 patients; 10.8% vs 3.2%; OR, 3.71; 95% CI, 1.49-9.2; P = .005) (Supplementary Table III, online only).

The absolute risk of stroke among asymptomatic women treated with CAS was 3%, with no significant differences compared with men (2.9%) (9 studies, 14,155 patients [5588 women, 8567 men]; OR, 1.09; 95% CI, 0.88-1.35; $\vec{r} = 7\%$; P = .42)^{9,11,15,18,51-54} (Fig 4). There was no sex difference in the absolute risk of death in asymptomatic patients treated with CAS (8 studies, 14,292 patients [5351 women, 8941 men]; OR, 1.16; 95% CI, 0.71-2.89; $\vec{r} = 19\%$; P = .55).^{9,11,18,20,51,53} However, this meta-analysis of observational studies showed that asymptomatic women treated with CAS had a significantly higher risk of in-hospital and

	Wom		Mer			Odds Ratio	Odds Ratio
tudy or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
1.1.1 Hospital, Asymptomatic							
Sisdas 2012 Rockman 2011 Seumenment	338	24521	284	24521	15.1%	1.19 [1.02, 1.40]	
tockman 2011, Asymptomati Jarac 2002, Asympotmatic	183 17	20584 787	261 16	27713	14.1× 3.7×	0.94 [0.78, 1.14] 2.17 [1.09, 4.31]	T
Subtotal (95% CI)	17	45892	10	53820	33.0%	1.16 [0.88, 1.54]	•
otal events	538		561				↓ ↓
leterogeneity: Tau ² = 0.04; Ch		df = 2 (F		r ² = 72%	6		
est for overall effect: Z = 1.07	(P = 0.29)	}					
4.1.2 1 month, Asymptomatic		2006	76	3666	10.28	1 49 11 09 2 021	
Calvillo—King 2010 CREST 2011, Howard	90 3	2996 191	75 5	3655 396	10.2%	1.48 [1.08, 2.02] 1.25 [0.30, 5.28]	
Javidovic 2015	22	530	22	1037	4.6%	2.00 [1.10, 3.64]	
Guzman 2014	2	120	3	245	0.7%	1.37 [0.23, 8.29]	
Hugi 2006	2	120	ō	252	0.2%	10.65 [0.51, 223.66]	
Im 2014	26	1718	39	2320	6.0%	0.90 [0.55, 1.48]	
Cuy 2014, Asymptomatic	474	86150	562	114737	16.2%	1.12 [0.99, 1.27]	•
ane 2003, Asymptomatic	0	58	0	142		Not estimable	
ee 2003, Asymptomatic	4	285	4	407	1.1%	1.43 [0.36, 5.78]	
ubke 2015, Asymptomatic	3	460	10	964	1.2%	0.63 [0.17, 2.29]	
Mattos 2001, Asymptomatic	04	154 332	2	247	0.2%	0.32 [0.02, 6.66]	
tockman 2001, Asymptomati Sternbach 2000	0	332 68	9	444 88	1.5%	0.59 [0.18, 1.93] 0.43 [0.02, 10.62]	
Subtotal (95% CI)	v	93182		124934	43.2%	1.19 [1.01, 1.40]	•
Fotal events	630		732				•
leterogeneity: Tau ² = 0.01; Ch		, df = 11		6); I ² = 9	×		
lest for overall effect: Z = 2.13							
122	-						
4.1.3 3 months, Asymptomat				A-4	*		
lugi 2006 Subtotal (95% CI)	3	120	2	252	0.7%	3.21 [0.53, 19.44]	
Subtotal (95% CI)	•	120	~	252	0.7%	3.21 [0.53, 19.44]	
Fotal events Heterogeneity: Not applicable	3		2				
Test for overall effect: $Z = 1.27$	(P = 0.21)						
		, ,					
4.1.4 4 years, Asymptomatic							
CREST 2011, Howard	5	191	6	396	1.6%	1.30 [0.42, 4.04]	
Subtotal (95% CI)		191		396	1.6%	1.30 [0.42, 4.04]	
Total events	5		6				
Heterogeneity: Not applicable	(n - 0.00						
rest for overall effect: Z = 0.46	(F = 0.05	,					
4.1.5 5 years, Asymptomatic							
ACST-2	25	541	54	1273	6.2%	1.09 [0.67, 1.78]	<u> </u>
CST 2004	12	539	18	1021	3.3%	1.27 [0.61, 2.65]	_
ane 2003, Asymptomatic	0	58	2	142	0.2%	0.48 [0.02, 10.16]	
Subtotal (95% CI)		1138		2436	9.8%	1.13 [0.75, 1.68]	+
Fotal events	37		74	.2			
Heterogeneity: $Tau^2 = 0.00$; Ch Fact for coursell effects $7 = 0.58$			· = 0.81);	r = 0%			
fest for overall effect: Z = 0.58	VL = 0.30	,					
4.1.6 10 years, Asymptomatic	:						
ACST-1 2010, Haliklay (1)	16	2658	53	5272	5.1%	0.60 [0.34, 1.05]	
CST-1 2010, Hallklay (2)	17	687	13	965	3.4%	1.86 [0.90, 3.85]	↓ • • •
Mattos 2001, Asymptomatic	9	154	38	247	3.2%	0.34 [0.16, 0.73]	
Subtotal (95% CI)		3499		6484	11.7%	0.72 [0.29, 1.79]	
fotal events	42		104				
Heterogeneity: $Tau^2 = 0.52$; Ch			(P = 0.00)	5); f = 6	18		
Test for overall effect: Z = 0.70	(P = 0.48)	,					
Fotal (95% CI)		144022		188322	100.0%	1.12 [0.96, 1.30]	L
Fotal events	1255		1461			[0.50, 1.50]	ľ
Heterogeneity: Tau ² = 0.03; Ch		. df = 22	-	2); $l^2 = 4$	2%		
Test for overall effect: Z = 1.47							0.02 0.1 1 1'0 5
lest for subgroup differences: ((P = 0.79)	$, ^2 = 0$	K.		Women Men
ootnotes		~~0					
$1) \leq 75$ years							
$(1) \ge 75$ years							

Fig 3. Stroke (A) and mortality (B) in men and women after endarterectomy of asymptomatic carotid artery stenosis. *Cl*, Confidence interval.

1-month MI than men (6 studies, 8445 patients; 1.2% vs 0.6%; OR, 2.01; 95% CI, 1.23-3.28; $l^2 = 0\%$; P = .005) (Fig 5; Supplementary Table IV [online only]).^{9,15,18,51-53}

The data on sex differences in the efficacy and safety of CAS in carotid stenosis comes from observational analyses of registries. Stroke and death among asymptomatic

В	Wom	en	Me	en		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	
4.3.1 Hospital, Asymptomatic						,	
Bisdas 2012	115	24521	118	24521	20.2%	0.97 [0.75, 1.26]	-
Rockman 2011, Asymptomati	77	20584	123	27713	16.5%	0.84 [0.63, 1.12]	
Sarac 2002, Asympotmatic	7	787	12	1586	1.5%	1.18 [0.46, 3.00]	_ _
Yavas 2010	0	16	1	57	0.1%	1.14 [0.04, 29.36]	
Subtotal (95% CI)		45908	_	53877	38.3%	0.92 [0.77, 1.11]	•
Total events	199		254				
Heterogeneity: Tau ² = 0.00; Ch	r ² = 0.84,	df = 3 (i)	= 0.84	; l ² = 0%			
Test for overall effect: Z = 0.84	$\langle P = 0.40 \rangle$)}					
4.3.2 1 month, Asymptomatic							
Guzman 2014	1	120	3	245	0.3%	0.68 [0.07, 6.59]	
Hugi 2006	0	120	3	252	0.2%	0.30 [0.02, 5.77]	
Jim 2014	11	1718	19	2320	2.4%	0.78 [0.37, 1.64]	
Kuy 2014, Asymptomatic	224	86150	356	114737	47.8%	0.64 [0.71, 0.99]	
Lee 2003, Asymptomatic	1	285	1	407	0.2%	1.43 [0.09, 22.95]	
Lubke 2015, Asymptomatic	0	460	2	964	0.1%	0.42 [0.02, 8.72]	
Sternbach 2000	0	68	0	88		Not estimable	
Subtotal (95% CI)		88921		119013	51.0%	0.83 [0.71, 0.98]	•
Total events	237		364				
Heterogeneity: Tau ² = 0.00; Ch Test for overall effect: Z = 2.24			P = 0.97)	; ² = 0%			
4.3.3 1 year, Asymptomatic							
Hugi 2006	1	120	12	252	0.3%	0.17 [0.02, 1.31]	
Subtotal (95% CI)		120		252	0.3%	0.17 [0.02, 1.31]	
Total events	1		12				
Heterogeneity: Not applicable Test for overall effect: Z = 1.70	(P = 0.09)}					
4.3.4 4 years, Asymptomatic							
Yavas 2010	1	16	6	53	0.3%	0.52 [0.06, 4.69]	
Subtotal (95% CI)		16	•	53	0.3%	0.52 [0.06, 4.69]	
Total events	1		6				
Heterogeneity: Not applicable Test for overall effect: $Z = 0.58$	(P = 0.56	i)					
4.3.5 5 years, Asymptomatic							
Kragsterman 2006	41	1737	80	3440	9.2%	1.02 [0.69, 1.49]	+
Lane 2003, Asymptomatic	4	58	9	58	0.9%	0.40 [0.12, 1.39]	
Subtotal (95% CI)		1795		3498	10.1%	0.78 [0.34, 1.77]	-
Total events	45		89				
Heterogeneity: Tau ² = 0.21; Ch Test for overall effect: Z = 0.60			P = 0.16)	; I ² = 497	6		
Total (95% CI)		136760		176693	100.0%	0.87 [0.78, 0.98]	•
Total events	483		745				
Heterogeneity: Tau ² = 0.00; Ch Test for overall effect: Z = 2.35 Test for subgroup differences: C	$\langle P = 0.02 \rangle$	2)					0.01 0.1 1 10 100 Women Men
Fig 3. Continued.							
-							

women treated with CAS were recorded in 2.6% to 5.4% of cases.^{9,15,51,55} In some of these studies, women were significantly more likely to develop stroke and death after CAS than men.^{11,52,53,56} In the study by Dua et al, female sex was associated with a high risk of postoperative stroke (OR, 12.59; 95% CI, 8.25-18.38; P < .001) and, together with CAS, it was one of the strongest risk factors for death (OR, 21.39; 95% CI, 5.49-33.39; P < .001).⁵² Other studies showed no between-sex differences in stroke and death in asymptomatic patients undergoing CAS.^{9,15,18,54} The risk of bias was acceptable.

DISCUSSION

We present data from our meta-analysis that collected evidence addressing revascularization of carotid artery stenosis in men and women covering the last 30 years of stroke evidence for this treatment. Although in some studies a higher perioperative risk with CAS and a higher stroke and death rate with CEA were reported, this did not result in a significant difference in the outcome after carotid revascularization in men and women, considering all endpoints.

Although there was a trend toward increased randomization of women over this period, women continue to be underrepresented in RCTs, and the percentage of women over 75 years of age are still low compared with that observed in the real clinical practice.⁵⁷ This under-enrollment was confirmed by a recent meta-analysis, underlining that this disparity persisted across all geographic regions, intervention types, and stroke types, apart from subarachnoid hemorrhage.⁵⁸

200
200
;

Fig 4. Stroke in men and women after stenting of asymptomatic carotid artery stenosis. *CI*, Confidence interval; *OR*, odds ratio.

	Wom		Mei			Odds Ratio	Odds Ratio	
Study or Subgroup		Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI	
3.5.1 Hospital, Asymptomat	ic							
Bisdas 2012	17	1773	5	1773	24.1%	3.42 [1.26, 9.30]		
Subtotal (95% CI)		1773		1773	24.1%	3.42 [1.26, 9.30]	-	
Total events	17		5					
Heterogeneity: Not applicable								
Test for overall effect: Z = 2.4	1 (P = 0)	.02)						
3.5.2 1 month, Asymptomat	tic							
CREST 2009, Howard	8	437	6	713	21.2%	2.20 [0.76, 6.38]		
CREST 2011, Howard	3	215	4	379	10.6%	1.33 [0.29, 5.98]		
De Rango 2010	2	306	1	778	4.2%	5.11 [0.46, 56.58]		
Jim 2014, Asymptomatic	9	743	12	1098	31.8%	1.11 [0.47, 2.65]		
Salinas-Aragón 2016, Asym	4	84	2	146	8.1%	3.60 [0.65, 20.09]	1	
Subtotal (95% CI)	-	1785	_	3114	75.9%	1.70 [0.97, 2.98]		
Total events	26		25				-	
Heterogeneity: Tau ² = 0.00; (9. df -	-).59): I	² = 0%			
Test for overall effect: Z = 1.6					•••-			
Total (95% CI)		3558		4887	100.0%	2.01 [1.23, 3.28]	•	
Total events	43		30				-	
Heterogeneity: $Tau^2 = 0.00$; (-	3. df -	5 (P = 1	0.52); 1	² = 0%		ahr alr da rh	
Test for overall effect: Z = 2.7							0.01 0.1 1 10	100
Test for subgroup differences			= 1 (P -	0.23)	P = 30 3	aw.	Women Men	

Fig 5. Myocardial infarction (MI) in men and women after stenting of asymptomatic carotid artery stenosis. *Cl.* Confidence interval.

Women are underrepresented in carotid revascularization trials, with the highest representation in the CREST-trial (35%). Apart from a higher rate of carotid atherosclerosis in men, the potential reasons for the underrepresentation of women in carotid revascularization trials may be the perceived technical difficulties (smaller internal carotid artery size in women) or a higher rate of peri- and postprocedural complications reported in women than in men.⁸

However, based on our data, we could not find a significant increased operative risk in women. In a consensus document published in 2013 on the management of women with carotid artery disease, bearing in mind the anatomical and technical differences and vascular and non-vascular comorbidities in men and women, a more tailored management for women was called for.⁵⁹ Current guidelines on the treatment of extracerebral vascular disease by the Society for Vascular Surgery do not give any sex-specific recommendation.⁶⁰ This applies also to the recently published guidelines on endarterectomy and stenting for carotid artery stenosis of the European Stroke Organisation. Sub-group analyses according to sex was performed, but due to the lack of interaction by sex for the main outcomes and low numbers of women included in RCTs, no specific recommendation for given.⁶¹ women was Considering the underenrollment of women, there could be potential risks of under-treatment, and it is important to state that, even with some studies reporting a higher perioperative risk in women, both sexes benefit likewise from revascularization. This was highlighted by a recently published algorithm for carotid stenosis in women.⁶² There are currently two ongoing trials comparing modern medical therapy with modern medical therapy and CAS/CEA in asymptomatic (CREST 2)⁶³ and in low-risk symptomatic patients (European Carotid Surgery Trial [ECST-2]).⁶⁴

A large RCT with a more pragmatic design, including an elderly population, may answer some questions about the risk and benefits of carotid intervention in women.⁶⁵

Limitations. Our systematic review is not without limitations. First, data are mostly based on cohort studies with possible inclusion bias. Although there are few RCTs in this systematic review, in these studies, patients were not randomized to men and women. Second, the authors were not contacted for the missing information and individual-based data of men and women due to the large number of studies included in this systematic review. Also, the management of carotid artery stenoses might have changed over time.

CONCLUSIONS

Overall, even considering the risk of bias, our data showed no significantly different outcomes in men and women after revascularization of symptomatic and asymptomatic carotid artery stenosis.

Further larger multicenter prospective research into these sex-specific differences is needed. More women have to be enrolled in RCTs, including women aged over 80 years, to better understand why these sex differences still exist and how we can tailor stroke treatment for both sexes.

AUTHOR CONTRIBUTIONS

Conception and design: CK, SL, YB, AL, ZG, VC

Analysis and interpretation: CK, YB, AL, MM, TK, DJ, JD, VC Data collection: CK, SL, YB, AL, CE, ZG, MM, TK, DJ, JD, VC Writing the article: CK, SL, ZG, VC

- Critical revision of the article: CK, SL, YB, AL, CE, ZG, MM, TK, DJ, JD, VC
- Final approval of the article: CK, SL, YB, AL, CE, ZG, MM, TK, DJ, JD, VC

Statistical analysis: AL

Obtained funding: Not applicable

Overall responsibility: CK

REFERENCES

- Petty GW, Brown RD Jr, Whisnant JP, Sicks JD, O'Fallon WM, Wiebers DO. Ischemic stroke subtypes: a population-based study of incidence and risk factors. Stroke 1999;30:2513-6.
- de Weerd M, Greving JP, Hedblad B, et al. Prevalence of asymptomatic carotid artery stenosis in the general population: an individual participant data meta-analysis. Stroke 2010;41:1294-7.
- Merz AA, Cheng S. Sex differences in cardiovascular ageing. Heart 2016;102:825-31.
- Schulz UG, Rothwell PM. Sex differences in carotid bifurcation anatomy and the distribution of atherosclerotic plaque. Stroke 2001;32:1525-31.
- Song JW, Cao Q, Siegler JE, Thon JM, Woo JH, Cucchiara BL. Sex differences in carotid plaque composition in patients with Embolic stroke of undetermined source. J Am Heart Assoc 2021;10:e020143.
- Krejza J, Arkuszewski M, Kasner SE, et al. Carotid artery diameter in men and women and the relation to body and neck size. Stroke 2006;37:1103-5.
- Carcel C, Woodward M, Wang X, Bushnell C, Sandset EC. Sex matters in stroke: a review of recent evidence on the differences between women and men. Front Neuroendocrinol 2020;59:100870.
- 8. Tsivgoulis G, Katsanos AH, Caso V. Under-representation of women in stroke randomized controlled trials: inadvertent selection bias leading to suboptimal conclusions. Ther Adv Neurol Disord 2017;10:241-4.
- Bisdas T, Egorova N, Moskowitz AJ, et al. The impact of gender on inhospital outcomes after carotid endarterectomy or stenting. Eur J Vasc Endovasc Surg 2012;44:244-50.
- Menyhei G, Bjorck M, Beiles B, et al. Outcome following carotid endarterectomy: lessons learned from a large international vascular registry. Eur J Vasc Endovasc Surg 2011;41:735-40.
- Rockman CB, Garg K, Jacobowitz GR, et al. Outcome of carotid artery interventions among female patients, 2004 to 2005. J Vasc Surg 2011;53:1457-64.
- Sarac TP, Hertzer NR, Mascha EJ, et al. Gender as a primary predictor of outcome after carotid endarterectomy. J Vasc Surg 2002;35: 748-53.
- Alamowitch S, Eliasziw M, Barnett HJ; North American Symptomatic Carotid Endarterectomy T, Group ASAT, Carotid Endarterectomy Trial G. The risk and benefit of endarterectomy in women with symptomatic internal carotid artery disease. Stroke 2005;36:27-31.
- Carotid Stenting Trialists C, Bonati LH, Dobson J, Algra A, et al. Shortterm outcome after stenting versus endarterectomy for symptomatic carotid stenosis: a preplanned meta-analysis of individual patient data. Lancet 2010;376:1062-73.
- Howard VJ, Lutsep HL, Mackey A, et al. Influence of sex on outcomes of stenting versus endarterectomy: a subgroup analysis of the Carotid Revascularization Endarterectomy versus Stenting Trial (CREST). Lancet Neurol 2011;10:530-7.
- Frawley JE, Hicks RG, Woodforth IJ. Risk factors for peri-operative stroke complicating carotid endarterectomy: selective analysis of a

prospective audit of 1000 consecutive operations. Aust N Z J Surg 2000;70:52-6.

- Guzman RP, Weighell W, Guzman C, Rodriguez-Leyva D. Female sex does not influence 30-day stroke and mortality rates after carotid endarterectomy. Ann Vasc Surg 2014;28:245-52.
- Jim J, Dillavou ED, Upchurch GR Jr, et al. Gender-specific 30-day outcomes after carotid endarterectomy and carotid artery stenting in the Society for Vascular Surgery Vascular Registry. J Vasc Surg 2014;59:742-8.
- 19. Kapral MK, Redelmeier DA. Carotid endarterectomy for women and men. J Womens Health Gend Based Med 2000;9:987-94.
- 20. Kuy S, Dua A, Desai SS, et al. Carotid endarterectomy national trends over a decade: does sex matter? Ann Vasc Surg 2014;28:887-92.
- 21. Lane JS, Shekherdimian S, Moore WS. Does female gender or hormone replacement therapy affect early or late outcome after carotid endarterectomy? J Vasc Surg 2003;37:568-74.
- Lee JW, Pomposelli F, Park KW. Association of sex with perioperative mortality and morbidity after carotid endarterectomy for asymptomatic carotid stenosis. J Cardiothorac Vasc Anesth 2003;17:10-6.
- Lubke T, Ahmad W, Koushk Jalali B, Brunkwall J. Gender-based 30day and long-term outcomes after carotid endarterectomy. Vasa 2015;44:289-95.
- 24. Mattos MA, Sumner DS, Bohannon WT, et al. Carotid endarterectomy in women: challenging the results from ACAS and NASCET. Ann Surg 2001;234:438-45; discussion: 445-6.
- Maxwell JG, Rutherford EJ, Covington DL, et al. Community hospital carotid endarterectomy in patients over age 75. Am J Surg 1990;160: 598-603.
- Riles TS, Imparato AM, Jacobowitz GR, et al. The cause of perioperative stroke after carotid endarterectomy. J Vasc Surg 1994;19:206-14; discussion: 215-6.
- Rockman CB, Svahn JK, Willis DJ, et al. Carotid endarterectomy in patients 55 years of age and younger. Ann Vasc Surg 2001;15:557-62.
- Schneider JR, Droste JS, Golan JF. Carotid endarterectomy in women versus men: patient characteristics and outcomes. J Vasc Surg 1997;25:890-6; discussion: 897-8.
- Tsivgoulis G, Krogias C, Georgiadis GS, et al. Safety of early endarterectomy in patients with symptomatic carotid artery stenosis: an international multicenter study. Eur J Neurol 2014;21. 1251-7, e75-e76.
- **30.** Weise J, Kuschke S, Bahr M. Gender-specific risk of perioperative complications in carotid endarterectomy patients with contralateral carotid artery stenosis or occlusion. J Neurol 2004;251:838-44.
- Cunningham EJ, Bond R, Mehta Z, et al. Long-term durability of carotid endarterectomy for symptomatic stenosis and risk factors for late postoperative stroke. Stroke 2002;33:2658-63.
- Paciaroni M, Eliasziw M, Kappelle LJ, Finan JW, Ferguson GG, Barnett HJ. Medical complications associated with carotid endarterectomy. North American symptomatic carotid endarterectomy trial (NASCET). Stroke 1999;30:1759-63.
- Chang JB, Stein TA. Ten-year outcome after saphenous vein patch angioplasty in males and females after carotid endarterectomy. Vasc Endovascular Surg 2002;36:21-7.
- 34. International Carotid Stenting Study i, Ederle J, Dobson J, Featherstone RL, et al. Carotid artery stenting compared with endarterectomy in patients with symptomatic carotid stenosis (International Carotid Stenting Study): an interim analysis of a randomised controlled trial. Lancet 2010;375:985-97.
- Brott TG, Hobson RW, Howard G, et al. Stenting versus endarterectomy for treatment of carotid-artery stenosis. N Engl J Med 2010;363: 11-23.
- 36. Eckstein HH, Ringleb P, Allenberg JR, et al. Results of the Stent-Protected Angioplasty versus Carotid Endarterectomy (SPACE) study to treat symptomatic stenoses at 2 years: a multinational, prospective, randomised trial. Lancet Neurol 2008;7:893-902.
- 37. Rothwell PM. ACST: which subgroups will benefit most from carotid endarterectomy? Lancet 2004;364:1122-3. author reply 5-6.
- Alamowitch S, Eliasziw M, Algra A, Meldrum H, Barnett HJ; Group NASCETN. Risk, causes, and prevention of ischaemic stroke in elderly patients with symptomatic internal-carotid-artery stenosis. Lancet 2001;357:1154-60.
- 39. Ederle J, Featherstone R, Brown M; Collaborators C. Long-term outcome of Endovascular treatment versus medical care for carotid

artery stenosis in patients not suitable for surgery and randomised in the carotid and Vertebral artery transluminal angioplasty study (CAVATAS). Cerebrovasc Dis 2009;28:1-7.

- 40. Ringleb P, Allenberg J, Berger J, et al. 30 day results from the SPACE trial of stent-protected angioplasty versus carotid endarterectomy in symptomatic patients: a randomised non-inferiority trial. Lancet 2006;368:1239-47.
- Halliday A, Harrison M, Hayter E, et al. 10-year stroke prevention after successful carotid endarterectomy for asymptomatic stenosis (ACST-1): a multicentre randomised trial. Lancet 2010;376:1074-84.
- 42. Halliday A, Mansfield A, Marro J, et al. Prevention of disabling and fatal strokes by successful carotid endarterectomy in patients without recent neurological symptoms: randomised controlled trial. Lancet 2004;363:1491-502.
- **43.** Davidovic L, Koncar I, Dragas M, et al. Female and obese patients might have higher risk from surgical repair of asymptomatic carotid artery stenosis. Ann Vasc Surg 2015;29:1286-92.
- Goldstein LB, McCrory DC, Landsman PB, et al. Multicenter review of preoperative risk factors for carotid endarterectomy in patients with ipsilateral symptoms. Stroke 1994;25:1116-21.
- 45. Hugl B, Oldenburg WA, Neuhauser B, Hakaim AG. Effect of age and gender on restenosis after carotid endarterectomy. Ann Vasc Surg 2006;20:602-8.
- Kragsterman B, Björck M, Lindbäck J, Bergqvist D, Pärsson H; (Swedvasc) SVR. Long-term survival after carotid endarterectomy for asymptomatic stenosis. Stroke 2006;37:2886-91.
- Sternbach Y, Perler BA. The influence of female gender on the outcome of carotid endarterectomy: a challenge to the ACAS findings. Surgery 2000;127:272-5.
- Yavas S, Mavioglu L, Kocabeyoglu S, et al. Is female gender really a risk factor for carotid endarterectomy? Ann Vasc Surg 2010;24:775-85.
- 49. Calvillo-King L, Xuan L, Zhang S, Tuhrim S, Halm EA. Predicting risk of perioperative death and stroke after carotid endarterectomy in asymptomatic patients: derivation and validation of a clinical risk score. Stroke 2010;41:2786-94.
- Halliday A, Bulbulia R, Bonati LH, et al. Second asymptomatic carotid surgery trial (ACST-2): a randomised comparison of carotid artery stenting versus carotid endarterectomy. Lancet 2021;398:1065-73.
- Werner N, Zeymer U, Mark B, et al. Carotid artery stenting in clinical practice: does sex matter? Results from the carotid artery stenting registry of the Arbeitsgemeinschaft Leitende Kardiologische Krankenhausärzte (ALKK). Clin Cardiol 2012;35:111-8.
- 52. Dua A, Romanelli M, Upchurch GR, et al. Predictors of poor outcome after carotid intervention. J Vasc Surg 2016;64:663-70.
- 53. Salinas-Aragon MA, Palacios-Rodríguez JM, García-Gutiérrez JC, et al. Impact of gender on short- and long-term morbidity and mortality after carotid stent angioplasty in a third-level hospital in Mexico. Rev Mex Cardiol 2016;27:34-43.
- 54. De Rango P, Parlani G, Caso V, et al. A comparative analysis of the outcomes of carotid stenting and carotid endarterectomy in women. J Vasc Surg 2010;51:337-44; discussion: 344.
- 55. Gray WA, Chaturvedi S, Verta P; Committees latE. Thirty-day outcomes for carotid artery stenting in 6320 patients from 2 prospective, multicenter, high-surgical-risk registries. Circ Cardiovasc Interv 2009;2:159-66.
- 56. Badheka AO, Chothani A, Panaich SS, et al. Impact of symptoms, gender, co-morbidities, and operator volume on outcome of carotid artery stenting (from the Nationwide Inpatient Sample [2006 to 2010]). Am J Cardiol 2014;114:933-41.
- Redon J, Olsen MH, Cooper RS, et al. Stroke mortality and trends from 1990 to 2006 in 39 countries from Europe and Central Asia: implications for control of high blood pressure. Eur Heart J 2011;32: 1424-31.
- Strong B, Pudar J, Thrift AG, et al. Sex disparities in Enrollment in recent randomized clinical trials of acute stroke: a meta-analysis. JAMA Neurol 2021;78:666-77.
- 59. De Rango P, Brown MM, Leys D, et al. Management of carotid stenosis in women: consensus document. Neurology 2013;80: 2258-68.
- AbuRahma AF, Avgerinos ED, Chang RW, et al. Society for Vascular Surgery clinical practice guidelines for management of extracranial cerebrovascular disease. J Vasc Surg 2022;75:4S-22S.

- Bonati LH, Kakkos S, Berkefeld J, et al. European Stroke Organisation guideline on endarterectomy and stenting for carotid artery stenosis. Eur Stroke J 2021;6:I-XLVII.
- 62. Rockman C, Caso V, Schneider PA. Carotid interventions for women: the hazards and benefits. Stroke 2022;53:611-23.
- **63.** Howard VJ, Meschia JF, Lal BK, et al. Carotid revascularization and medical management for asymptomatic carotid stenosis: protocol of the CREST-2 clinical trials. Int J Stroke 2017;12:770-8.
- 64. Cheng SF, van Velzen TJ, Gregson J, et al. The 2nd European Carotid Surgery Trial (ECST-2): rationale and protocol for a randomised clinical trial comparing immediate revascularisation versus optimised medical therapy alone in patients with symptomatic and

asymptomatic carotid stenosis at low to intermediate risk of stroke. Trials 2022;23:606.

65. Bereznyakova O, Dewar B, Dowlatshahi D, et al. Benefit of carotid revascularisation for women with symptomatic carotid stenosis: protocol for a systematic review. BMJ Open 2019;9:e032140.

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Additional material for this article may be found online at www.jvascsurg.org.

SUPPLEMENTARY APPENDIX 1 (online only). Methods: Supplemental Information

Medical Subject Headings (MeSH) for the Search Used. 'Carotid stenosis' OR 'carotid artery stenosis' OR 'carotid artery obstruction' AND 'carotid artery surgery' OR ('carotid artery' AND 'surgery') OR 'angioplasty' OR 'stent*' OR 'angioplasty, balloon' OR 'percutaneous transluminal angioplasty' OR 'endarterectomy" AND 'treatment outcome' OR 'postoperative complications' OR 'myocardial infarction' OR 'heart infarction' OR 'stroke' OR 'brain ischemia' OR 'cerebrovascular accident' OR 'death' OR 'death, sudden, cardiac' OR 'mortality' OR 'sudden death' AND 'females' OR 'males' OR 'women' OR 'men' OR 'gender difference' OR 'sex difference' OR 'sex factor*' OR 'gender factor'.

Inclusion and Exclusion Criteria for the Search Used. Symptomatic carotid artery stenosis.

Women and men with symptomatic carotid artery stenosis; Carotid endarterectomy in women; Carotid

endarterectomy in men; Stroke, hemorrhage, mortality.

Women and men with symptomatic carotid artery stenosis; Carotid stenting in women; Carotid stenting in men; Stroke, hemorrhage, mortality.

Asymptomatic carotid artery stenosis.

Women and men with asymptomatic carotid artery stenosis; Carotid endarterectomy in women; Carotid endarterectomy in men; Stroke, hemorrhage, mortality.

Women and men with asymptomatic carotid artery stenosis; Carotid stenting in women; Carotid stenting in men; Stroke, hemorrhage, mortality.

Exclusion Criteria. Patients: Women and men without carotid artery stenosis; Did not evaluate carotid endarterectomy or carotid stenting; Did not study Stroke, hemorrhage, mortality. Study designs such as reviews, letter to editor, case report, commentary, or editorial.

SUPPLEMENTARY APPENDIX 2 (online only).

PRISMA 2020 Checklist

		-	
	ltem		Location where item is
Section and Topic	#	Checklist item	reported
Title			
Title	1	Identify the report as a systematic review.	Title page
Abstract			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	Abstract
Introduction			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	р. 3
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	p. 3-4
Methods			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	Suppl.File 1
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	p. 4
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	p.4 Suppl.File1
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	p.4

Continued.

	Item		Location where item is
Section and Topic	#	Checklist item	reported
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	p.4
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (eg for all measures, time points analyses), and if not, the methods used to decide which results to collect.	
	10b	List and define all other variables for which data were sought (eg participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	p.4 t
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	p.4

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Continued.			
	ltem		Location where item is
Section and Topic	#	Checklist item	reported
Effect measures	12	Specify for each outcome the effect measure(s) (eg risk ratio, mean difference) used in the synthesis or presentation of results.	p.4
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (eg tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	p.4
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	p.4
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	p.4
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta- analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	p.4
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (eg subgroup analysis, meta-regression).	n.a.

Continued.

			Location
Section and Topic	ltem #	Checklist item	where item is reported
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	p. 4
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	p.4 and figures
Results			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	
Study characteristics	17	Cite each included study and present its characteristics.	p. 11
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	p.5 cont.
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (eg confidence/credible interval), ideally using structured tables or plots.	p.5 cont.

(Continued)

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Continued.

	ltem		Location where item is
Section and Topic	#	Checklist item	reported
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	p.5 cont.
	20b	Present results of all statistical syntheses conducted. If meta- analysis was done, present for each the summary estimate and its precision (eg confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	Figs 1-5 and suppl.
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	p. 5 cont.
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	p.5 cont.
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	p. 5 cont.
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	p. 5 cont.
Discussion			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	p.8 cont.
	23b	Discuss any limitations of the evidence included in the review.	p.8
	23c	Discuss any limitations of the review processes used.	p.9
			(Continued)

Continued.

Continued.			
Section and Topic	ltem #	Checklist item	Location where item is reported
	23d	Discuss implications of the results for practice, policy, and future research.	p.9
Other information			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	Not registered
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	p.4
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	n/a
Support	25	Describe sources of financial or non- financial support for the review, and the role of the funders or sponsors in the review	p.10
Competing interests	26	Declare any competing interests of review authors.	p. 10
Availability of data, code, and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	p.5 cont.
		IE, Bossuyt PM, et al. The eline for reporting systemat	

2021; 372:n71. For more information, visit: http://www.prisma-statement.org/. Supplementary Table I (online only). Vascular events, length of stay (LOS), and complications, in men and women after endarterectomy of symptomatic carotid artery stenosis

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	P value
Stroke						
Hospital	2.1	1.9	5 (54,585)	1.11 [0.89-1.38]	57%; .05	.36
1 month	5.0	4.5	19 (37,333)	1.10 [1.00-1.22]	0%; .88	.05
1 year	0.0	1.9	1 (93)	0.43 [0.02-10.89]	NA	.61
2 years	5.3	6.5	2 (1015)	0.99 [0.40-2.42]	47%; .17	.98
4 years	8.8	5.9	2 (926)	1.57 [0.81-3.05]	37%; .21	.18
5 years	14.1	13.1	2 (1597)	1.18 [0.71-1.94]	13%; .28	.53
10 years	11.4	12.4	3 (3946)	0.81 [0.55-1.19]	67%; .05	.28
TIA						
Hospital	0.6	0.3	1 (1049)	1.91 [0.27-13.62]	NA	.52
1 month	1.9	1.4	3 (1449)	1.64 [0.71-3.80]	0%; .95	.25
2 years	3.9	4.8	1 (426)	0.80 [0.30-2.15]	NA	.66
Death						
Hospital	1.2	1.6	4 (8036)	0.83 [0.57-1.21]	0%; .52	.33
1 month	1.0	0.8	10 (76,604)	1.15 [0.94-1.39]	10%; .35	.17
1 year	NR	NR	NR	NR	NR	NR
2 years	NR	NR	NR	NR	NR	NR
4 years	8.3	9.2	1 (122)	0.90 [0.18-4.46]	NA	.9
5 years	14.8	16.8	4 (1786)	0.84 [0.64-1.09]	0%; .53	.19
10 years	27.1	37.8	1 (615)	0.61 [0.43-0.87]	NA	.006
Stroke or death						
Hospital	4.4	3.9	2 (5417)	1.09 [0.84-1.43]	0%; .84	.51
1 month	9.4	8.9	13 (14,360)	1.08 [0.93-1.27]	8%; .37	.3
4 months	7.2	5.0	2 (2565)	1.49 [1.04-2.12]	0%; .34	.03
3 years	NR	NR	NR	NR	NR	NR
4 years	6.2	5.4	1 (653)	1.16 [0.58-2.30]	NA	.67
5 years	12.4	8.9	3 (2331)	1.47 [0.94-2.29]	59%; .09	.09
MI						
Hospital	1.3	1.0	1 (4368)	1.34 [0.76-2.36]	NA	.32
1 month	1.5	1.4	4 (24,284)	0.98 [0.61-1.56]	33%; .21	.92
Stroke, MI or death						
Hospital	8.2	8.6	1 (697)	0.95 [0.54-1.65]	NA	.85
1 month	5.7	5.0	3 (3928)	1.14 [0.67-1.95]	59%; .09	.62
4 years	7.5	7.7	1 (653)	0.97 [0.53-1.78]	NA	.92
LOS, hospital, days						
Overall	6.4 ± 11.8	5.8 ± 11.7	2 (21,177)	0.52 [0.21-0.83]	0%; .55	.001
Restenosis						
1 month	3.6	3.2	1 (212)	1.12 [0.21-5.94]	NA	.9
1 year	NR	NR	NR	NR	NR	NR
5 years	11.4	3.3	1 (615)	3.79 [1.89-7.61]	NA	.0002
10 years	17.2	6.7	1 (615)	2.81 [1.66-4.75]	NA	.0001
Reintervention						
1 month	3.2	2.4	1 (811)	1.32 [0.56-3.10]	NA	.52

Supplementary Table I (online only) Continued.

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	P value
CHF						
1 month	0.6	0.6	1 (811)	1.05 [0.17-6.32]	NA	.96
Cranial nerve palsy						
Overall	8.2	4.3	1 (821)	1.98 [1.08-3.64]	NA	.03
Hematoma						
Overall	8.2	5.2	1 (821)	1.64 [0.91-2.95]	NA	.1
CHF, Congestive heart fail	lure; CI, confidence ir	nterval; I ² , heterog	geneity; ICU, intensiv	e care unit; <i>MI</i> , myocardial in	farction; <i>n</i> , numbe	er of studies; N,

number of patients; *NA*, not applicable; *NE*, not estimable; *OR*, odds ratio; *P*, statistical significance value.

Supplementary Table II (online only). Vascular events, length of stay (LOS), and complications in men and women after stenting of symptomatic carotid artery stenosis

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	<i>P</i> value
Stroke						
Hospital	7.5	7.1	3 (939)	1.16 [0.59-2.30]	34%; .22	.67
1 month	7.0	8.5	5 (2436)	0.83 [0.59-1.16]	0%; .60	.28
2 years	8.2	9.6	1 (607)	0.84 [0.44-1.57]	NA	.58
4 years	9.6	5.8	1 (668)	1.71 [0.95-3.08]	NA	.08
TIA						
1 month	4.5	5.3	1 (255)	0.83 [0.22-3.13]	NA	.79
Death						
Hospital	1.9	1.6	3 (3421)	0.88 [0.43-1.77]	38%; .20	.71
1 month	2.2	2.0	5 (3984)	1.20 [0.61-2.35]	37%; .17	.60
Stroke or death						
Hospital	5.3	4.8	2 (2961)	0.97 [0.64-1.48]	21%; .26	.9
1 month	7.7	6.9	6 (3661)	1.15 [0.87-1.51]	0%; .61	.33
4 months	8.5	9.0	1 (1725)	0.93 [0.64-1.36]	NA	.72
4 years	9.6	5.8	1 (668)	1.71 [0.95-3.08]	NA	.08
MI						
Hospital	1.7	2.6	1 (466)	0.66 [0.18-2.37]	NA	.53
1 month	1.7	4.5	5 (2980)	0.59 [0.05-6.74]	88%, <0.001	.67
Stroke, MI, or death						
1 month	7.8	7.8	4 (2725)	1.07 [0.67-1.70]	42%; .16	.79
4 months	7.9	8.7	1 (853)	0.91 [0.53-1.56]	NA	.73
LOS, hospital, days						
Overall	6.7 ± 1.4	5.4 ± 1.2	2 (721)	-0.09 [-0.27 to 0.08]	0%; .80	.29

CI, Confidence interval: *I*², heterogeneity: *ICU*, intensive care unit; *MI*, myocardial infarction; *n*, number of studies; *N*, number of patients; *NA*, not applicable; *NE*, not estimable; *OR*, odds ratio; *P*, statistical significance value.

Supplementary Table III (online only). Vascular events, length of stay (LOS), and complications in men and women after endarterectomy of asymptomatic carotid artery stenosis

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	P value
Stroke						
Hospital	1.2	1.0	3 (99,712)	1.16 [0.88-1.54]	72%; .03	.29
1 month	0.7	0.6	12 (218,116)	1.19 [1.01-1.40]	9%; .36	.03
3 months	2.5	0.8	1 (372)	3.21 [0.53-19.44]	NA	.21
1 year	NR	NR	NR	NR	NR	NR
2 years	NR	NR	NR	NR	NR	NR
4 years	2.6	2.0	1 (587)	1.30 [0.42-4.04]	NA	.65
5 years	3.3	3.0	3 (3574)	1.13 [0.75-1.68]	0%; .81	.56
10 years	1.2	1.6	3 (9983)	0.72 [0.29-1.79]	81%; .005	.48
TIA						
Hospital	0.9	0.6	1 (2373)	1.57 [0.58-4.24]	NA	.37
1 month	0.6	0.1	3 (1220)	3.56 [0.46-27.67]	0%; .34	.23
3 months	2.5	0.0	1 (372)	15.04 [0.77-293.58]	NA	.07
2 years						
Death						
Hospital	0.4	0.5	4 (99,785)	0.92 [0.77-1.11]	0%; .84	.4
1 month	0.27	0.32	7 (207,934)	0.83 [0.71-0.98]	0%; .97	.03
3 months	NR	NR	NR	NR	NR	NR
l year	0.8	4.8	1 (372)	0.17 [0.02-1.31]	NA	.09
2 years	NR	NR	NR	NR	NR	NR
4 years	6.3	11.3	1 (69)	0.52 [0.06-4.69]	NA	.56
5 years	2.5	2.5	2 (5293)	0.78 [0.34-1.77]	49%; .16	.55
10 years	NR	NR	NR	NR	NR	NR
Stroke or death						
Hospital	1.7	1.6	2 (51,415)	1.35 [0.68-2.71]	82%; .02	.39
1 month	3.2	2.1	5 (10,218)	1.44 [1.13-1.85]	0%; .53	.004
4 months	NR	NR	NR	NR	NR	NR
3 years	2.2	1.8	1 (1560)	1.27 [0.61-2.65]	NA	.53
4 years	2.6	2.0	1 (587)	1.30 [0.42-4.04]	NA	.65
5 years	5.4	4.0	1 (1560)	1.36 [0.83-2.21]	NA	.22
MI						
Hospital	0.8	0.5	1 (49,042)	1.48 [1.17-1.85]	NA	.0008
1 month	0.90	0.85	5 (206,360)	1.06 [0.96-1.16]	0%; .72	.23
Stroke, MI or death						
Hospital	5.3	1.6	1 (463)	3.43 [1.10-10.69]	NA	.03
1 month	3.1	3.2	2 (4625)	0.96 [0.69-1.34]	0%; .86	.81
LOS, hospital, days						
Overall	2.6±16.0	2.3±16.0	2 (201,579)	0.24 [0.10-0.38]	0%; .87	.0006
Restenosis						
1 month	NR	NR	NR	NR	NR	NR
l year	10.8	3.2	1 (372)	3.71 [1.49-9.20]	NA	.005
5 years	NR	NR	NR	NR	NR	NR
10 years	NR	NR	NR	NR	NR	NR
Reintervention						
1 month	2.4	2.9	3 (1264)	0.80 [0.39-1.67]	0%; .74	.56

Supplementary Table III (online only) Continued.

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	P value
CHF						
1 month	1.7	0.8	2 (848)	2.07 [0.58-7.42]	0%; .95	.26
Arrhythmia						
1 month	0.0	1.1	1 (156)	0.43 [0.02-10.62]	NA	.6
Cranial nerve palsy						
Overall	2.5	2.8	1 (372)	0.90 [0.23-3.53]	NA	.88
Hematoma						
Overall	1.7	2.0	1 (372)	0.84 [0.16-4.38]	NA	.83
Wound infections						
Overall	0.8	0.8	1 (372)	1.05 [0.09-11.70]	NA	.97
CHF, Congestive heart fail	lure; CI, confidence	interval; I ² , hetero	ogeneity; <i>ICU</i> , intensi	ve care unit; <i>MI</i> , myocardial infa	arction; <i>n</i> , number	of studies; <i>N</i> ,

number of patients; NA, not applicable; NE, not estimable; OR, odds ratio; P, statistical significance value.

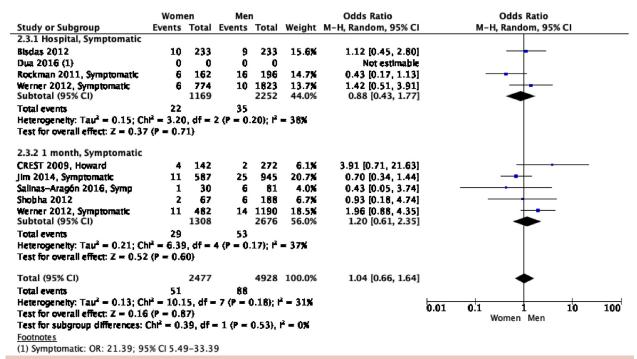
Supplementary Table IV (online only). Vascular events, length of stay (LOS), and complications in men and women after stenting of asymptomatic carotid artery stenosis

	Incidence, %					
Outcome	Women	Men	n (N)	OR [95% CI]	l ² ; <i>P</i>	<i>P</i> value
Stroke						
Hospital	2.1	1.9	2 (6279)	1.13 [0.79-1.62]	0%; .78	.49
1 month	3.5	3.2	5 (5471)	1.04 [0.63-1.74]	53%; .08	.87
2 years	NR	NR	NR	NR	NR	NR
4 years	5.0	4.7	2 (2405)	1.10 [0.74-1.64]	0%; .93	.64
TIA						
1 month	3.3	3.7	1 (1084)	0.87 [0.42-1.81]	NA	.71
Death						
Hospital	0.7	0.6	3 (8404)	1.09 [0.61-1.95]	0%; .47	.78
1 month	1.1	0.9	5 (5888)	1.12 [0.43-2.86]	49%; .12	.82
Stroke or death						
Hospital	2.5	2.5	2 (5600)	0.99 [0.69-1.42]	0%; .34	.95
1 month	2.7	3.3	5 (7479)	0.84 [0.60-1.17]	15%; .32	.3
4 years	4.2	4.0	1 (594)	1.06 [0.46-2.47]	NA	.89
MI						
Hospital	1.0	0.3	1 (3546)	3.42 [1.26-9.30]	NA	.02
1 month	1.5	0.8	5 (4899)	1.70 [0.97-2.98]	0%; .59	.07
Stroke, MI, or death						
1 month	5.7	4.1	4 (3815)	1.46 [0.95-2.24]	36%; .19	.09
LOS, hospital, days						
Overall	3.1 ± 3.0	2.7 ± 3.0	1 (3546)	0.40 [0.20-0.60]	NA	.29
Hematoma						
1 month	1.6	1.3	1 (1084)	1.28 [0.43-3.76]	NA	.66

CI. Confidence interval; l^2 , heterogeneity; *ICU*, intensive care unit; *MI*, myocardial infarction; *n*, number of studies; *N*, number of patients; *NA*, not applicable; *NE*, not estimable; *NR*, not reported; *OR*, odds ratio; *P*, statistical significance value.

Study or Subarows	Wom		Me		Wainht	Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
2.2.1 Hospital, Symptomatic				210-	F 9-4	A CA 14 37 4 341	
Bisdas 2012	16	2184	23	2164	5.3%	0.69 [0.37, 1.32]	
Rockman 2011, Symptomatic	22	1037	39	1449	7.0%	0.78 [0.46, 1.33]	
Sarac 2002, Sympotmatic	7	361	6	666	2.4%	1.68 [0.60, 4.67]	
řavas 2010 Subtotal (95% CI)	0	26 3608	3	107 4428	0.3X 15.0%	0.56 [0.03, 11.24] 0.83 [0.57, 1.21]	•
Fotal events	45		73				
Heterogeneity: $Tau^2 = 0.00$; Cl Test for overall effect: $Z = 0.91$			(P = 0.5	2); I ² = (0%		
2.2.2 1 month, Symptomatic							
Alamowitch 2005	17	753	14	1810	4.5%	2.96 [1.45, 6.04]	
Guzman 2014	2	243	3	438	0.8%	1.20 [0.20, 7.25]	
Im 2014	12	960	14	1494	3.9%	1.34 [0.62, 2.91]	_
Kapral 2000	4	506	9	1140	1.9%	1.00 [0.31, 3.27]	
Kuy 2014, Symptomatic	158	8254	208	12112	16.6%	1.12 [0.91, 1.36]	
ee 2003, Symptomatic	Ō	315	2	496	0.3%	0.31 [0.02, 6.55]	
Lubke 2015, Symptomatic	ĩ	128	2	328	0.5%	1.28 [0.12, 14.28]	
Menyhel 2011 < 75 Y	32	9017		20561	9.5%	0.94 [0.62, 1.41]	
Menyhel 2011, > 75 Y	36	6123	-	11500	9.7%	1.01 [0.67, 1.51]	
Schneider 1997	1	155	0	271	0.3%	5.27 [0.21, 130.20]	
Subtotal (95% CI)	-	26454	v	50150		1.15 [0.94, 1.39]	•
Total events	263		397				•
Heterogeneity: $Tau^2 = 0.01$; Cl Test for overall effect: $Z = 1.3$!			(P = 0.3	5); f² = 1	10%		
2.2.3 4 years, Symptomatic							
řavas 2010 Subtotal (95% CI)	2	24 24	9	96 98	1.1% 1.1%	0.90 [0.16, 4.46] 0.90 [0.18, 4.46]	
Fotal events	2		9				
Heterogeneity: Not applicable Fest for overall effect: Z = 0.13) (P = 0.9	0)					
2.2.4 5 year, Symptomatic							
Chang 2002	43	247	67	366	9.3%	0.95 [0.62, 1.44]	_
Kragsterman 2006	39	229	62	402	9.3%	0.80 [0.53, 1.22]	+
ane 2003, Symptomatic	14	57	14	57	3.3%	1.00 [0.43, 2.35]	_
Schneider 1997	6	155	22	271	2.9%	0.46 [0.18, 1.15]	<u> </u>
Subtotal (95% CI)	-	688		1098	24.8%	0.84 [0.64, 1.09]	•
Fotal events	102		185				-
Heterogeneity: $Tau^2 = 0.00$; Cl Test for overall effect: $Z = 1.31$			(P = 0.5	3); I ² = (0%		
2.2.5 10 year, Symptomatic							
Chang 2002	67	247	139	368	11.3%	0.61 [0.43, 0.87]	
Subtotal (95% CI)		247		368	11.3%	0.61 [0.43, 0.87]	◆
	67		139				-
Fotal events							
l otal events Heterogeneity: Not applicable Test for overall effect: Z = 2.73	3 (P = 0.0)	06)					
Heterogeneity: Not applicable	3 (P = 0.0	31021		56142	100.0%	0.95 [0.80, 1.12]	•
Heterogeneity: Not applicable Test for overall effect: $Z = 2.7$	3 (P = 0.04 479		603	56142	100.0%	0.95 [0.80, 1.12]	4
Heterogeneity: Not applicable Test for overall effect: Z = 2.7; Total (95% Cl)	479	31021					0.01 0.1 1 10 1

Supplementary Fig 1 (online only). Mortality in men and women after endarterectomy of symptomatic carotid artery stenosis. *Cl*, Confidence interval.



Supplementary Fig 2 (online only). Mortality in men and women after stenting of symptomatic carotid artery stenosis. *Cl*, Confidence interval; *OR*, odds ratio.