

LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE



Doig, D; Turner, EL; Dobson, J; Featherstone, RL; de Borst, GJ; Brown, MM; Richards, T; the ICSS Investigators (2014) Incidence, Impact, and Predictors of Cranial Nerve Palsy and Haematoma Following Carotid Endarterectomy in the International Carotid Stenting Study. *European journal of vascular and endovascular surgery* . ISSN 1078-5884 DOI: 10.1016/j.ejvs.2014.08.002

Downloaded from: <http://researchonline.lshtm.ac.uk/2017944/>

DOI: [10.1016/j.ejvs.2014.08.002](https://doi.org/10.1016/j.ejvs.2014.08.002)

#### Usage Guidelines

Please refer to usage guidelines at <http://researchonline.lshtm.ac.uk/policies.html> or alternatively contact [researchonline@lshtm.ac.uk](mailto:researchonline@lshtm.ac.uk).

Available under license: <http://creativecommons.org/licenses/by/2.5/>

# Incidence, Impact, and Predictors of Cranial Nerve Palsy and Haematoma Following Carotid Endarterectomy in the International Carotid Stenting Study

D. Doig<sup>a</sup>, E.L. Turner<sup>b</sup>, J. Dobson<sup>c</sup>, R.L. Featherstone<sup>a</sup>, G.J. de Borst<sup>d</sup>, M.M. Brown<sup>a,\*</sup>, T. Richards<sup>e</sup>, on behalf of the ICSS Investigators<sup>f</sup>

<sup>a</sup>Institute of Neurology, University College London, London, UK

<sup>b</sup>Department of Biostatistics and Bioinformatics and Duke Global Health Institute, Duke University, Durham, NC, USA

<sup>c</sup>Department of Medical Statistics, London School of Hygiene and Tropical Medicine, London, UK

<sup>d</sup>Department of Vascular Surgery, University Medical Centre Utrecht, Utrecht, The Netherlands

<sup>e</sup>Department of Surgical and Interventional Sciences, University College London, London, UK

## WHAT THIS PAPER ADDS

Perioperative cranial nerve palsy remains an important complication of carotid endarterectomy, and occurred in 5.5% of patients in a recent large randomized trial. There is a significant association between the occurrence of perioperative haematoma and perioperative cranial nerve palsy. Patients can be reassured that many cranial nerve palsies following carotid endarterectomy are transient and non-disabling. Women should be warned about an increased risk of cranial nerve palsy.

**Objective:** Cranial nerve palsy (CNP) and neck haematoma are complications of carotid endarterectomy (CEA). The effects of patient factors and surgical technique were analysed on the risk, and impact on disability, of CNP or haematoma in the surgical arm of the International Carotid Stenting Study (ICSS), a randomized controlled clinical trial of stenting versus CEA in patients with symptomatic carotid stenosis.

**Materials and methods:** A per-protocol analysis of early outcome in patients receiving CEA in ICSS is reported. Haematoma was defined by the surgeon. CNP was confirmed by an independent neurologist. Factors associated with the risk of CNP and haematoma were investigated in a binomial regression analysis.

**Results:** Of the patients undergoing CEA, 45/821 (5.5%) developed CNP, one of which was disabling (modified Rankin score = 3 at 1 month). Twenty-eight (3.4%) developed severe haematoma. Twelve patients with haematoma also had CNP, a significant association ( $p < .01$ ). Independent risk factors modifying the risk of CNP were cardiac failure (risk ratio [RR] 2.66, 95% CI 1.11 to 6.40), female sex (RR 1.80, 95% CI 1.02 to 3.20), the degree of contralateral carotid stenosis, and time from randomization to treatment >14 days (RR 3.33, 95% CI 1.05 to 10.57). The risk of haematoma was increased in women, by the prescription of anticoagulant drugs pre-procedure and in patients with atrial fibrillation, and was decreased in patients in whom a shunt was used and in those with a higher baseline cholesterol level.

**Conclusions:** CNP remains relatively common after CEA, but is rarely disabling. Women should be warned about an increased risk. Attention to haemostasis might reduce the incidence of CNP. ICSS is a registered clinical trial: ISRCTN 25337470.

© 2014 Published by Elsevier Ltd on behalf of European Society for Vascular Surgery. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/3.0/>).

Article history: Received 4 April 2014, Accepted 1 August 2014, Available online 2 October 2014

**Keywords:** Carotid atherosclerosis, Carotid artery stenosis, Carotid endarterectomy, Cranial nerves, Haematoma

## INTRODUCTION

Evidence from randomized trials of medical therapy versus carotid endarterectomy (CEA) for symptomatic stenosis of the carotid artery<sup>1</sup> has led to the recommendation that CEA should be performed in patients with symptomatic carotid artery stenosis to reduce the long-term risk of recurrent stroke or TIA.<sup>2</sup> The combined rate of stroke or death at 30 days following CEA in NASCET, ECST, and the VA trials was 7.1% (95% CI 6.3 to 8.1%).<sup>3</sup> However, the primary endpoints of these trials did not include cranial nerve palsy (CNP) or haematoma.

<sup>f</sup> For a list of ICSS investigators see *Lancet* 2010;375:985–997.

\* Corresponding author. M.M. Brown, Box 6, National Hospital for Neurology and Neurosurgery, Queen Square, London, WC1N 3BG, UK.

E-mail address: [martin.brown@ucl.ac.uk](mailto:martin.brown@ucl.ac.uk) (M.M. Brown).

1078-5884/© 2014 Published by Elsevier Ltd on behalf of European Society for Vascular Surgery. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/3.0/>).

<http://dx.doi.org/10.1016/j.ejvs.2014.08.002>

Although less extensively studied, the surgical complications of CNP and haematoma have long been recognized following CEA,<sup>4</sup> and have been associated with an increased risk of stroke or death.<sup>5</sup> Nerves affected include the mandibular branch of the facial nerve, vagal, glossopharyngeal, hypoglossal, and accessory nerves,<sup>4,6,7</sup> and therefore CNP has the potential to cause significant postoperative morbidity.

Carotid angioplasty and stenting (CAS) was developed as an alternative to CEA, in part to avoid these hazards of a surgical incision. However, the results of recent large randomized trials, including the International Carotid Stenting Study (ICSS),<sup>8</sup> have consistently shown that CAS carries a higher risk of non-disabling stroke than CEA within 30 days of the procedure, with no significant difference in the rates of disabling stroke or death.<sup>9</sup> ICSS was a randomized controlled multicentre open clinical trial that randomized patients with symptomatic carotid stenosis to CEA or CAS. In this study, the incidence and severity of CNP and haematoma in ICSS, and risk factors for their development, were studied, to identify groups of patients at higher risk and determine whether these complications merit consideration in selection of a revascularization procedure.

## METHODS

### Study design

**Patient selection and protocol design.** The protocol for ICSS is published elsewhere.<sup>10</sup> Patients over 40 years old were eligible for randomization in ICSS if they had more than 50% recently symptomatic carotid stenosis suitable for either CAS or CEA, and were clinically stable. Patients were excluded if they had a major stroke with poor recovery of function, if their vascular anatomy rendered CAS or CEA unsuitable, if the stenosis was caused by non-atheromatous disease, if cardiac bypass was planned within 1 month of the revascularization procedure, or if there had been previous revascularization of the symptomatic artery.

Carotid endarterectomy in ICSS was performed according to the surgeon's usual practice: local, general, or combined anaesthesia was allowed for the procedure. The type of arterial reconstruction to be carried out was not specified in the protocol, nor was the choice of peri-procedural medication.

**Outcome events.** Technical details of the surgical procedure and the occurrence of cranial nerve palsy or haematoma were

reported by trial investigators. All patients were then re-assessed at 1 month after the procedure by a neurologist or investigator under their supervision. CNPs were adjudicated internally at the ICSS trial office and judged to be disabling if the patient's score on the modified Rankin Scale (mRS) increased to 3 or more at 30 days after the procedure, where that increase was attributable to the CNP. Investigators were additionally asked to complete a questionnaire (Appendix I) giving details of the clinical consequences of CNP and whether or not the lesion resolved during subsequent follow-up in the trial. Haematoma was classified as severe if it required re-operation, transfusion, or prolonged hospital stay.

**Statistical analysis.** The data were analysed per-protocol; only patients in whom the randomly allocated procedure was initiated were included in this analysis. Patients who crossed over or received CEA after an attempt at stenting were excluded. A procedure was deemed to have been initiated if the patient underwent either local or general anaesthesia prior to commencement of surgery. Risk factors for CNP and haematoma were examined sequentially in a univariable binomial regression analysis using maximum likelihood estimation. The risk ratio for each factor was estimated with a 95% confidence interval. Wald tests were used for continuous and binary predictors, with an overall likelihood ratio test for categorical predictors of more than two levels. A multivariable model was developed using a forward stepwise based approach. Patients with missing data were excluded from each relevant analysis. Analyses were performed with Stata (StataCorp. 2011. *Stata Statistical Software: Release 12*. College Station, TX: StataCorp LP).

## RESULTS

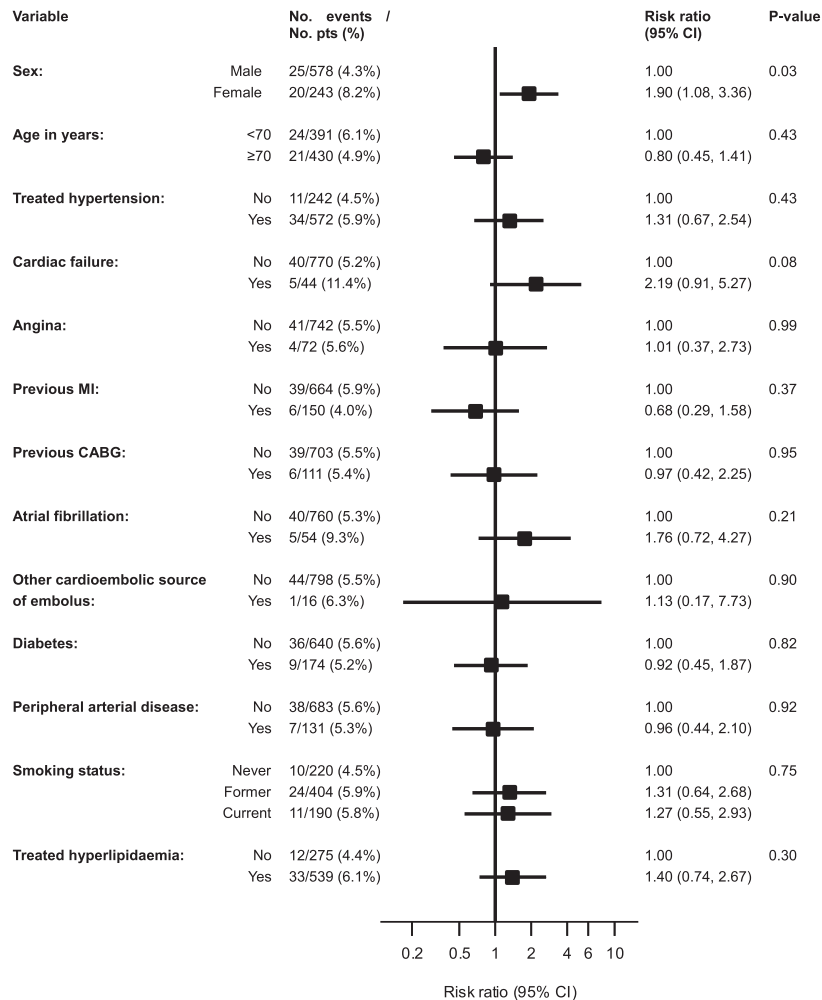
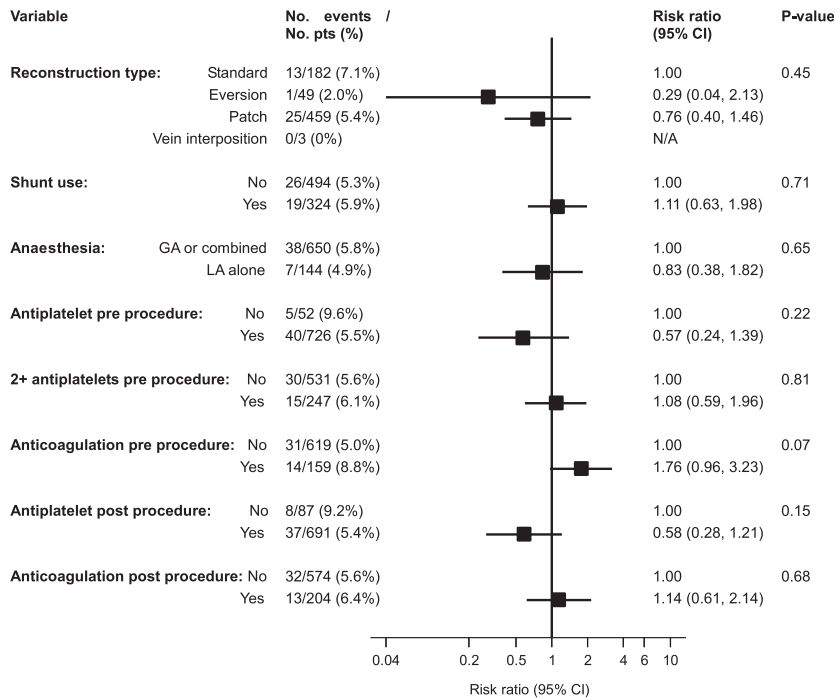
### Cranial nerve palsy

In ICSS, 1713 patients were randomized. Of 858 patients randomized to CEA, the allocated procedure was initiated in 821 (95.7%). Four of 821 patients (0.5%) died between initiation of the procedure and 30 days post-procedure. Forty-five of 821 patients (5.5% of initiated CEAs) were reported to have CNP within 30 days of the procedure. The results of adjudication of which cranial nerves were affected are presented in Table 1. A total of 50 CNPs were reported: facial ( $n = 23$ ), vagus (6), hypoglossal (13), glossopharyngeal (4), accessory (1), and trigeminal (1); in

**Table 1.** Summary of cranial nerve palsies (CNPs) within 30 days of endarterectomy in ICSS per-protocol participants in whom the procedure was initiated ( $n = 821$ ).

Cranial nerve	Number of CNPs ( $n = 50$ in 45 patients)	Number of disabling CNPs (mRS $\geq 3$ because of CNP)	Number of CNPs confirmed persisting after 30 days
Facial	23	0	4
Hypoglossal	13	0	2
Vagus	6	0	4
Accessory	1	0	0
Glossopharyngeal	4	1	1
Trigeminal	1	0	0
Undetermined	2	0	0

CNP = cranial nerve palsy; ICSS = International Carotid Stenting Study; mRS = modified Rankin Scale score.



**Figure 1.** Univariable predictors of risk of cranial nerve palsy within 30 days of endarterectomy in ICSS per-protocol participants in whom the procedure was initiated (n=821).

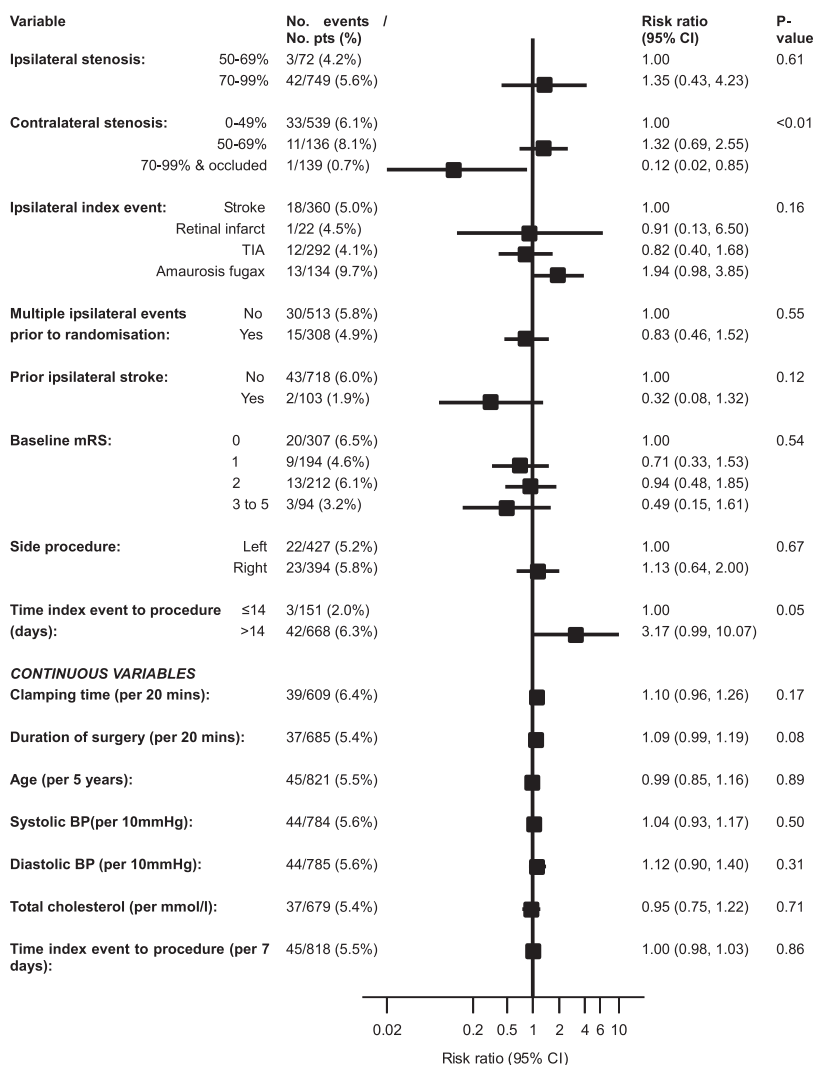


Figure 1. (continued).

two patients it was not possible to determine which cranial nerve was affected. One CNP was judged to be disabling with an mRS of 3 at 1-month follow-up. This patient had glossopharyngeal nerve palsy with impairment in swallowing requiring placement of a naso-gastric feeding tube. Two cranial nerve palsies were reported in each of five trial participants. In those patients with CNP where symptomatic resolution was confirmed ( $n = 20$ ), the median duration of symptoms before resolution was 30 days (minimum 2 days, maximum 520 days). The exact duration of symptoms in the remainder was undetermined. In only two of

821 patients (0.2%) were the symptoms reported to have not resolved during follow-up of 6.4 and 3.1 years, respectively. One of these patients experienced voice hoarseness caused by vocal cord paresis following vagal nerve injury. The other experienced uplifting of the mouth on one side as a result of facial nerve injury.

The results of the risk factor analysis are presented in Fig. 1. Statistically significant predictors of CNP in univariable analysis were female sex (risk ratio [RR] 1.90, 95% CI 1.08 to 3.36,  $p = .03$ ) and a high degree of contralateral carotid artery stenosis. Other demographic and technical

Table 2. Independent predictors of risk of cranial nerve palsy within 30 days of carotid endarterectomy in ICSS per-protocol participants ( $n = 805$ ) in whom the procedure was initiated. Patients with missing data were excluded from this analysis.

Variable	Adjusted risk ratio (95% CI)	Adjusted p value
Cardiac failure	2.66 (1.11 to 6.40)	.03
Female sex	1.80 (1.02 to 3.20)	.04
Time from randomization to treatment >14 days	3.33 (1.05 to 10.57)	.04
Degree of contralateral stenosis		Overall $p < .01$
0–50%	1.00	
50–69%	1.18 (0.62 to 2.27)	.62
>70%	0.13 (0.02 to 0.91)	.04

**Table 3.** Numbers (%) of patients with haematoma by presence of cranial nerve palsy (CNP)<sup>a</sup> within 30 days of carotid endarterectomy in ICSS per-protocol participants (n=821).

		Haematoma		
		Number (%)	Yes (%)	Total (%)
CNP	No (%)	738 (95.1)	38 (4.9)	776 (100)
	Yes (%)	33 (73.3)	12 (26.7)	45 (100)
	Total	771	50	821

<sup>a</sup>  $p < .01$  by Fisher's exact test of independence between haematoma and CNP.

factors, including the type of arterial reconstruction, type of anaesthesia or shunt use did not predict CNP. Independent predictors of CNP in multivariable analysis, summarised in Table 2, were cardiac failure (RR 2.66, 95% CI 1.11 to 6.40,  $p = .03$ ), female sex (RR 1.80, 95% CI 1.02 to 3.20,  $p = .04$ ), the degree of contralateral carotid stenosis, and time to operation of >14 days after the day of randomization (RR 3.33, 95% CI 1.05 to 10.57,  $p = .04$ ).

### Haematoma

Of 821 patients in whom the surgical procedure was initiated, 50 (6.1%) developed neck haematoma. Twenty-eight of the 821 (3.4%) were classified as severe.

The results of univariable regression analysis for the risk factors for haematoma development are presented in Appendix II. Statistically significant predictors of increased risk of haematoma were: anticoagulant prescription pre-operatively (RR 1.83, 95% CI 1.04 to 3.23,  $p = .04$ ), previous cardiac bypass graft surgery (CABG) (RR 2.46, 95% CI 1.37 to 4.42,  $p < .01$ ), atrial fibrillation (RR 2.29, 95% CI 1.08 to 4.85,  $p = .03$ ), and the duration of arterial clamping in minutes (RR per each extra 20 minutes 1.13, 95% CI 1.04 to 1.24,  $p < .01$ ). Factors associated with a decreased risk of postoperative haematoma were shunt use (RR 0.54, 95% CI 0.29 to 0.99,  $p = .05$ ), antiplatelet agent prescription prior to the procedure (RR 0.44, 95% CI 0.21 to 0.93,  $p = .03$ ) and each 1 mmol/l increase in cholesterol at baseline (RR 0.69, 95% CI 0.55 to 0.88,  $p < .01$ ). Other demographic and technical factors did not predict haematoma.

The results of multivariable analysis of predictors of risk for haematoma are presented in Appendix III. Independent predictors of increased risk were being female (RR 2.03, 95% CI 1.13 to 3.62,  $p = .02$ ), having atrial fibrillation (RR 2.38, 95% CI 1.07 to 5.27,  $p = .03$ ), and the prescription of anticoagulant pre-procedure (RR 1.86, 95% CI 1.01 to 3.42,  $p = .05$ ). Independent factors reducing the risk of

haematoma were shunt use (RR 0.40, 95% CI 0.21 to 0.80,  $p < .01$ ) and each 1 mmol/l increase in the patient's baseline cholesterol level (RR 0.68, 95% CI 0.54 to 0.86,  $p < .01$ ).

Twelve of 45 (26.7%) patients with CNP also suffered haematoma, versus 38/776 (4/9%) of patients without CNP. There was a significant association between these complications as detailed in Table 3 ( $p < .01$ , Fisher's exact test).

### Impact on trial outcomes

Table 4 details the impact of adding CNP to the combined incidence of stroke, myocardial infarction (MI), or death in ICSS in a post-hoc analysis comparing CEA with CAS. There was no significant difference in the combined risk of stroke, MI, death, or CNP, nor was there a significant difference in the incidence of disabling stroke, disabling CNP, or death between the two trial arms.

### DISCUSSION

In ICSS, CNP developed in 5.5% of ICSS patients undergoing CEA, and haematoma in 6.1%. There was a statistically significant association between the two outcomes. The largest available series of patients studied pre- and post-operatively, in the European Carotid Surgery Trial (ECST), found a motor CNP rate of 5.1% and a long-term CNP rate of 0.5% at 4 months.<sup>11</sup> Likewise, NASCET reported an overall risk of postoperative CNP of 8.6%, of which the majority were mild in severity,<sup>12</sup> suggesting that CNP rates remain constant over time.

Patients should be made aware of these common complications and the likely clinical effects, including sensory and possible motor consequences. They can be reassured from the evidence presented here that postoperative CNP is rarely disabling (risk around 1 in 1000 operations), but should be warned that the symptoms of the CNP may persist for several weeks or longer.

There are several other findings of interest from this study: to the authors' knowledge, the association between female sex and CNP has not previously been described, and this finding is worth confirming in another cohort of patients. One possible explanation for higher risk in female patients may be more challenging surgical anatomy and the smaller average diameter of the carotid artery.<sup>13</sup> Female patients were also at higher risk of haematoma. Combined with other reports of a higher perioperative stroke risk in

**Table 4.** Composite outcome events within 30 days of carotid stenting (CAS) versus carotid endarterectomy (CEA) in ICSS, with or without the addition of cranial nerve palsy (CNP), in per-protocol participants.

Endpoint	CAS (n = 828) No. events (%)	CEA (n = 821) No. events (%)	Risk ratio (95% CI)	Risk difference (95% CI)	p (chi-square)
Stroke, MI or death <sup>a</sup>	61 (7.4%)	33 (4.0%)	1.83 (1.21, 2.77)	3.3% (1.1, 5.6)	<.01
Stroke, MI, death or CNP	62 (7.5%)	76 (9.3%)	0.81 (0.59, 1.12)	-1.8% (-4.4, 0.9)	.20
Disabling stroke or death	26 (3.1%)	18 (2.2%)	1.43 (0.79, 2.59)	0.9% (-0.6, 2.5)	.23
Disabling stroke, disabling CNP or death	27 (3.3%)	19 (2.3%)	1.41 (0.79, 2.51)	0.9% (-0.6, 2.5)	.24

<sup>a</sup> Reported in *Lancet* 2010;376:985-997.

symptomatic women undergoing CEA versus men,<sup>14</sup> and evidence that the net benefit of CEA in women is lower than in men,<sup>15</sup> an increased incidence of CNP should be borne in mind when advising female patients of their risk of complications following CEA.

Other risk factors for CNP identified in this analysis, including the degree of contralateral carotid stenosis, are statistically significant predictors but are more difficult to link clinically with the outcome and have not, to the authors' knowledge, been reported by other groups.

The association of neck haematoma with pre-operative anticoagulation is not surprising, but emphasizes the importance of careful surgical technique to mitigate the risk of haematoma in anticoagulated patients. A risk of haematoma in ICSS of 6.1% is similar to the rate of "wound complications" seen in large case series<sup>16</sup> and a severe haematoma risk of 3.4% is similar to the reported risk of re-exploration of the surgical wound required in patients undergoing CEA while on antiplatelet medication.<sup>17</sup> However, although there is concern about perioperative bleeding in patients on dual antiplatelet therapy, the incidence of haematoma was actually decreased in this study in patients taking these medications, perhaps because surgeons took more care with haemostasis in these patients.

Some systematic reviews have included CNP in a composite outcome event of death or neurological complications up to 30 days after treatment.<sup>18</sup> ICSS compared CAS with CEA on the assumption that if CAS could avoid neck incision, CNP, and haematoma with no excess risk of stroke, then it could provide a beneficial alternative to CEA for the prevention of recurrent stroke in patients with symptomatic carotid stenosis.<sup>10</sup> In ICSS, as reported here, the total number of events in the composite cluster of any stroke, MI, death, or CNP was greater after CEA compared with CAS, but the numbers in the cluster of disabling stroke, disabling CNP or death were greater after CAS than after CEA. Neither difference was statistically significant. It is concluded, therefore, that it would not be appropriate to base treatment considerations concerning the choice of CAS versus CEA on the basis of composite short-term endpoints including CNP.

### Limitations of the analysis

The present analysis has some important limitations. In some patients, information regarding baseline risk factors was unavailable, and information about the duration of CNP symptoms was limited. Multiple comparisons without statistical correction raise the possibility of obtaining a Type I (false positive) error. Results could be confirmed in other patient cohorts. This is not a randomized comparison of surgical techniques or perioperative processes of care, and it is possible that unmeasured confounders are associated with the risk of CNP or haematoma.

### Conclusion

CNP remains a relatively common complication of CEA, but in many patients is transient. Haematoma is similarly common, and there is a statistical association between

haematoma and CNP. Women should be warned about an increased risk of CNP. Scrupulous attention to haemostasis might reduce the incidence of CNP. Fortunately, prolonged disability or permanent symptoms as a result of haematoma or CNP are rare, and thus, in the authors' opinion, do not warrant inclusion in composite endpoints for future trials of carotid revascularization, but nevertheless one in 821 CEA patients in ICSS had permanent impairment of swallowing caused by cranial nerve palsy.

### ACKNOWLEDGEMENTS

DD and RLF collated the data. RLF is the trial manager of ICSS. DD wrote the first draft of the manuscript. ELT and JD performed statistical analyses. ELT completed this work at the Department of Medical Statistics, London School of Hygiene and Tropical Medicine. TR and GJdB designed this study and edited the manuscript. MMB is the chief investigator of ICSS. All authors were involved in reviewing the manuscript prior to submission.

### CONFLICT OF INTEREST

None.

### FUNDING

ICSS was funded by the UK Medical Research Council (MRC) and managed by the UK National Institute for Health Research (NIHR) on behalf of the MRC-NIHR partnership (grant numbers G0300411 and EME 09/800/14 respectively). The views and opinions expressed herein are those of the authors and do not necessarily reflect those of the NIHR Health Services Research programme of the Department of Health. Additional funding was supplied by grants from the Stroke Association (grant numbers TSA2005/01 and TSA2007/12), Sanofi-Synthelabo and the European Union. MMB's Chair in Stroke Medicine is supported by the Reta Lila Weston Trust for Medical Research. DD and RLF were supported by a grant from the Medical Research Council. This work was undertaken at University College London, which received a proportion of funding from the UK Department of Health's National Institute for Health Research Biomedical Research Centres funding scheme. The funders and sponsors of the study had no role in study design, data collection, data analysis, data interpretation or the writing of this paper.

### APPENDIX A. SUPPLEMENTARY DATA

Supplementary data related to this article can be found online at <http://dx.doi.org/10.1016/j.ejvs.2014.08.002>.

### REFERENCES

- 1 Rerkasem K, Rothwell PM. Systematic review of randomized controlled trials of carotid endarterectomy for symptomatic carotid stenosis. *Stroke* 2011;**42**:e543–4.
- 2 Liapis CD, Sir Bell PRF, Mikhailidis D, Sivenius J, Nicolaidis A, Fernandes e Fernandes J, et al. on behalf of the ESVS Guidelines Collaborators. ESVS guidelines. Invasive treatment of carotid stenosis: indications, techniques. *Eur J Vasc Endovasc Surg* 2009;**37**:S1–19.

- 3 Rothwell PM, Eliasziw M, Gutnikov SA, Fox AJ, Taylor DW, Mayber MR, et al. for the Carotid Endarterectomy Trialists' Collaboration. Analysis of pooled data from the randomized controlled trials of endarterectomy for symptomatic carotid stenosis. *Lancet* 2003;**361**:107–16.
- 4 Massey EW, Heyman A, Utley C, Haynes C, Fuchs J. Cranial nerve paralysis following carotid endarterectomy. *Stroke* 1984;**15**:157–9.
- 5 Greenstein AJ, Chassin MR, Wang J, Rockman CB, Riles TS, Tuhim S, et al. Association between minor and major surgical complications after carotid endarterectomy: results of the New York Carotid Artery Surgery Study. *J Vasc Surg* 2007;**46**:1138–46.
- 6 Maniglia AJ, Han PD. Cranial nerve injuries following carotid endarterectomy: an analysis of 336 procedures. *Head Neck* 1991;**13**:121–4.
- 7 Ballotta E, Da Giau G, Renon L, Narne S, Saladini M, Abbruzzese E, et al. Cranial and cervical nerve injuries after carotid endarterectomy: a prospective study. *Surgery* 1999;**125**:85–91.
- 8 International Carotid Stenting Study Investigators. 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.
- 9 Carotid Stenting Trialists' Collaboration. Short-term outcome after stenting versus endarterectomy for symptomatic carotid stenosis: a preplanned meta-analysis of individual patient data. *Lancet* 2010;**376**:1062–73.
- 10 Featherstone RL, Brown MM, Coward LJ. International Carotid Stenting Study: protocol for a randomised clinical trial comparing carotid stenting with endarterectomy in symptomatic carotid artery stenosis. *Cerebrovasc Dis* 2004;**18**:69–74.
- 11 Cunningham EJ, Bond R, Mayberg MR, Warlow CP, Rothwell PM. Risk of persistent cranial nerve injury after carotid endarterectomy. *J Neurosurg* 2004;**101**:445–8.
- 12 Ferguson GG, Eliasziw M, Barr HWK, Clagett GP, Barnes RW, Wallace MC, et al. The North American Symptomatic Carotid Endarterectomy Trial: surgical results in 1415 patients. *Stroke* 1999;**30**:1751–8.
- 13 Krejza J, Arkuszewski M, Kasner SE, Weigle J, Ustymowicz A, Hurst RW, et al. Carotid artery diameter in men and women and relation to body and neck size. *Stroke* 2006;**37**:1103–5.
- 14 Rockman CB, Garg K, Jacobowitz GR, Berger JS, Mussa FF, Cayne NS, et al. Outcome of carotid artery interventions among female patients, 2004 to 2005. *J Vasc Surg* 2011;**53**:1457–64.
- 15 Rothwell PM, Eliasziw M, Gutnikov SA, Warlow CP, Harnett HJM. for the Carotid Endarterectomy Trialists' Collaboration. Endarterectomy for symptomatic carotid stenosis in relation to clinical subgroups and timing of surgery. *Lancet* 2004;**363**:915–24.
- 16 Press MJ, Chassin MR, Wang J, Tuhim S, Halm EA. Predicting medical and surgical complications of carotid endarterectomy: comparing the risk indexes. *Arch Intern Med* 2006;**166**:914–20.
- 17 Oldag A, Schreiber S, Schreiber S, Heinze H-J, Meyer F, Weber M, et al. Risk of wound haematoma at carotid endarterectomy under dual antiplatelet therapy. *Langenbecks Arch Surg* 2012;**397**:1275–82.
- 18 Bonati LH, Lyrer P, Ederle J, Featherstone R, Brown MM. Percutaneous transluminal balloon angioplasty and stenting for carotid artery stenosis. *Cochrane Database Syst Rev* 2012;**9**:CD000515. <http://dx.doi.org/10.1002/14651858.CD000515.pub4>.