Open versus endoscopic carpal tunnel release: A systematic review and meta-analysis of randomized controlled trials

CURRENT STATUS: UNDER REVIEW

Yueying Li
China-Japan Union Hospital of Jilin University

Wenqi Luo
China-Japan Union Hospital of Jilin University

Guangzhi Wu
China-Japan Union Hospital of Jilin University

Shusen Cui
China-Japan Union Hospital of Jilin University

Zhan Zhang
China-Japan Union Hospital of Jilin University

Xiaossong Gu
China-Japan Union Hospital of Jilin University

Corresponding Author
2622489738@qq.com
ORCiD: https://orcid.org/0000-0002-8973-9201

DOI:
10.21203/rs.2.22968/v2

SUBJECT AREAS
Orthopedics

KEYWORDS
Carpal tunnel syndrome; Complications; Endoscopic carpal tunnel release; Meta-analysis; Open carpal tunnel release; Randomized controlled trial
Abstract

**Background:** Endoscopic carpal tunnel release (ECTR) and open carpal tunnel release (OCTR) both have advantages and disadvantages for the treatment of carpal tunnel syndrome (CTS). We compared the effectiveness and safety of ECTR and OCTR based on evidence from a high-level randomized controlled trial.

**Methods:** We comprehensively searched PubMed, EMBASE, Cochrane Library, Web of Science, and Medline to identify relevant articles published until August 2019. Data regarding operative time, grip strength, Boston Carpal Tunnel Questionnaire scores, digital sensation, patient satisfaction, key pinch strength, return to work time, and complications were extracted and compared. All mean differences (MD) and odds ratios (OR) were expressed as ECTR relative to OCTR.

**Results:** Twenty-eight studies were included in our meta-analysis. ECTR was associated with significantly higher satisfaction rates (MD, 3.13; 95% confidence interval [CI], 1.43 to 4.82; P = 0.0003), greater key pinch strengths (MD, 0.79 kg; 95% CI, 0.27 to 1.32; P = 0.003), earlier return to work times (MD, -7.25 days; 95% CI, -14.31 to -0.19; P = 0.04), higher transient nerve injury rates (OR, 4.87; 95% CI, 1.37 to 17.25; P = 0.01), and a lower incidence of scar-related complications (OR, 0.20; 95% CI, 0.07 to 0.59; P = 0.004). There were no significant differences between the two methods in terms of permanent nerve injury (OR, 1.93; 95% CI, 0.58 to 6.40; P = 0.28).

**Conclusions:** Overall, evidence from randomized controlled trials indicates that ECTR results in better recovery of daily life functions than OCTR, as revealed by higher satisfaction rates, greater key pinch strengths, earlier return to work times, and fewer scar-related complications. Our findings suggest that patients with CTS can be effectively managed with ECTR.

1. **Background**

Carpal tunnel syndrome (CTS), known as compressive median mononeuropathy at the wrist, causes tingling, numbness, and pain along the radial side of the hand [1]. The reported estimates for its annual prevalence range from 0.18% to 5% [2–5]. CTS can be treated surgically or non-surgically; however, non-surgical management that involves wrist splinting, corticosteroid injections, and physiotherapy, is preferred over surgical management for mild and moderate CTS [6,7]. Surgical
treatments for CTS, including the open carpal tunnel release (OCTR) and endoscopic carpal tunnel release (ECTR) approach, are generally reserved for patients with severe symptoms or those who experienced conservative treatment failure [8,9].

OCTR is a well-established surgical treatment for CTS [10]. However, it is associated with potential complications such as persistent weakness, pillar pain, formation of hypertrophic scars in the incisions that cross the wrist, scar tenderness, slow recovery, and a higher incidence of persistent pain [11]. In an attempt to avoid these complications, Chow [12] and Okutsu et al. [13] first reported the use of ECTR for the treatment of CTS in the English literature in 1989. This method allows for smaller skin incisions and better esthetic results than OCTR [1,14,15]. Nevertheless, ECTR is technically difficult, time consuming, and associated with incomplete transverse carpal ligament release and neurovascular injury [16–20]. Several meta-analyses have compared various measures of efficacy and safety between ECTR and OCTR [15,21–23]. However, these investigations failed to separate subgroups according to different follow-up times and utilized limited evaluations of patient outcomes; therefore, it is not clear which approach is associated with better clinical results [24,25]. Therefore, we performed a meta-analysis of published results to compare the effectiveness and safety between ECTR and OCTR according to randomized controlled trial (RCT) evidence. Specifically, we sought to determine if ECTR was superior to OCTR in terms of patient satisfaction, functional recovery, and complications.

2. Methods

2.1. Literature search

The meta-analysis was performed according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines [26]. Two authors independently used PubMed, EMBASE, Cochrane Library, Web of Science, and Medline databases to search for relevant publications.

Publications from the inception of each database to August 10, 2019 were searched. The keywords used in the searches were “carpal tunnel” plus “open incision” and “carpal tunnel” plus “endoscopic.” To identify other relevant studies, we manually scanned the reference lists of the relevant articles that were discovered using these search terms.
2.2. Criteria for selected trials

A study was included if it was an RCT that compared OCTR and ECTR. The exclusion criteria were as follows: 1) descriptive or graphic outcomes with no standard deviation values, 2) studies that evaluated revision surgery, 3) studies that did not report the follow-up interval, 4) studies that reported only limited qualitative findings, 5) studies published in a language other than English or Chinese, and 6) abstracts, laboratory or anatomic studies, review or technique articles, commentaries, and nontherapeutic studies. Finally, two investigators independently reviewed all selected studies for inclusion.

2.3. Data extraction

Two independent reviewers extracted data from the included studies. Any discrepancy in data interpretation was resolved either by discussion or by involving a third reviewer until a consensus was reached for all items.

The data extracted from eligible studies included publication year, country of origin, sample size, intervention details, follow-up interval, and outcomes. If outcome data and measures of variance were reported graphically but were omitted from the body of the text, plot-digitizing software (Plot Digitizer Version 2.6.4; Joseph Huwaldt and Scott Steinhorst, http://www.plot-digitizer.com-about.com/) was used to quantify these data. The pooled analysis outcome parameters were as follows: operation duration; scores on several clinical indexes, including the Boston Carpal Tunnel Questionnaire Symptom Severity Scale (BCTQ-S), Boston Carpal Tunnel Questionnaire Functional Status Scale (BCTQ-F), Two-point Discrimination test, and Semmes-Weinstein monofilament test; grip strength; key pinch strength; time to return to work (RTW); patients’ subjective ratings of their satisfaction with symptom improvement following CTS release based on a scale of 0 to 100 points; and postoperative complications.

2.4. Quality assessment

The level of evidence was assessed according to the Grading of Recommendations Assessment, Development, and Evaluation (GRADE) guidelines [27]. At least two reviewers independently assessed the risk of bias, and disagreements were resolved through discussion.
2.5. Statistical analysis
Continuous data were analyzed using the inverse-variance statistical method and computation of the mean difference (MD) and 95% confidence interval (CI). Dichotomous data were analyzed using the Mantel-Haenszel statistical method and computation of the odds ratio (OR) and 95% CI. All MD and OR values were calculated using the results from OCTR as the reference values. In addition, \( \chi^2 \) and \( I^2 \) tests were used to assess statistical heterogeneity. Significant heterogeneity was indicated when the P value from the \( \chi^2 \) test was < 0.10 or when the \( I^2 \) value exceeded 50%. If heterogeneity was present, a random-effects model was applied to assess the pooled result of the outcome measure; otherwise, a fixed-effects model was used. All tests were two-tailed, and the threshold for statistical significance was set at P < 0.05. The funnel plot method and Egger’s test were utilized to evaluate the publication bias. The data were further analyzed using Review Manager (version 5.3; The Nordic Cochrane Centre, Cochrane Collaboration, Copenhagen, Denmark).

3. Results
3.1. Selected studies and characteristics
Figure 1 summarizes the number of articles identified, reviewed, and included in the final analysis. A total of 5,654 relevant articles were initially identified from PubMed (n = 1,416), EMBASE (n = 1,755), Cochrane Library (n = 248), Web of Science (n = 1,130), Medline (n = 1,105), and reference lists (n = 0). After exclusion of duplicates, 2,248 articles remained. Reviews of the titles and abstracts reduced the number of articles to 103, and a more detailed review reduced this number further to 28 articles for final inclusion in the meta-analysis. Twenty-seven articles were in English and one was in Chinese. The characteristics of the included articles are summarized in Table 1.

3.2. Quality assessment
According to the GRADE guidelines, 19 RCTs reported adequate methods for random sequence generation. Only 8 RCTs had low risks of detection bias for outcomes. The majority of RCTs (25/28) failed to report the blinding status of patients, study personnel, and outcome assessors. Attrition bias was judged as low risk for 22 RCTs. All RCTs were at a low risk of selective reporting of outcomes (Fig. 2).
3.3. Meta-analysis results

There were no significant differences in the operative time (MD, -5.81 min; 95% CI, -17.85 to 6.23; P = 0.34; n = 261; random-effects model, with a heterogeneity of $I^2 = 99\%$; $P < 0.00001$; Fig. 3) [28-31], grip strength at 3 months post-surgery (MD, 1.99 kg; 95% CI, -0.43 to 4.42; $P = 0.11$; n = 297; fixed-effects model, with a heterogeneity of $I^2 = 0\%$; $P = 0.79$; Fig. 4) [9,32], BCTQ-S score at 1 year post-surgery (MD, 0.15; 95% CI, -0.04 to 0.35; $P = 0.13$; n = 592; random-effects model, with a heterogeneity of $I^2 = 92\%$; $P < 0.00001$; Fig. 5) [25,33,34], and BCTQ-F score at 1 year post-surgery (MD, 0.17; 95% CI, -0.02 to 0.36; $P = 0.08$; n = 592; random-effects model, with a heterogeneity of $I^2 = 91\%$; $P < 0.00001$; Fig. 6) [25,33,34] between the ECTR and OCTR groups. Similarly, there were no differences in digital sensation, including the Semmes-Weinstein monofilament test score at 3 months post-surgery (MD, 0.06; 95% CI, -0.09 to 0.21; $P = 0.43$; n = 297; fixed-effects model, with a heterogeneity of $I^2 = 0\%$; $P = 0.65$; Fig. 7) [9,32] and Two-point Discrimination test score at 1 year post-surgery (MD, -0.16; 95% CI, -0.45 to 0.12; $P = 0.26$; n = 402; fixed-effects model, with a heterogeneity of $I^2 = 35\%$; $P = 0.20$; Fig. 8) [30,34,35], between the two groups.

3.3.1. Satisfaction rate

The overall level of satisfaction with the outcome was based on a scale of 0 to 100 points. Two articles provided comparative data on the satisfaction rate [32,34]. A portion of the data from Zhang et al. [34] reported a satisfaction rate of up to 90%, with high heterogeneity; therefore, some of the satisfaction data from that study were excluded from the present meta-analysis. The pooled data of the two articles showed that the satisfaction rate in the ECTR group was significantly higher than that in the OCTR group (MD, 3.13; 95% CI, 1.43 to 4.82; $P = 0.0003$; n = 303; fixed-effects model, with a heterogeneity of $I^2 = 0\%$; $P = 0.57$) [32,34], and the clinical heterogeneity $I^2$ was null (Fig. 9).

3.3.2. Key pinch strength

The pooled data showed that the key pinch strength of patients who were treated with ECTR was significantly greater than the key pinch strength of patients who were treated with OCTR at 3-months post-surgery (MD, 0.79 kg; 95% CI, 0.27 to 1.32; $P = 0.003$; n = 297; fixed-effects model, with a
heterogeneity of $I^2 = 0\%; P = 0.70$ [9,32] (Fig. 10).

3.3.3. RTW

Four studies [31,36-38] evaluated the time needed to return to work for patients who underwent CTS. The pooled data showed that the RTW times were significantly faster in patients in the ECTR group than those in the OCTR group (MD, -7.25 days; 95% CI, -14.31 to -0.19; $P = 0.04$; $n = 357$; random-effects model, with a heterogeneity of $I^2 = 98\%; P < 0.00001$) (Fig. 11); however, divergences between studies resulted in large between-study heterogeneity.

3.3.4. Complications

Twenty-five studies [9,24,28-50] included complete complication rate data and were included in the pooled analysis of overall complications. There were no significant differences between all complications rates (OR, 1.06; 95% CI, 0.69 to 1.64; $P = 0.78$; $n = 2320$; fixed-effects model, with a heterogeneity of $I^2 = 16\%; P = 0.27$) (Fig. 12). The rates of transient nerve injury were higher in patients who underwent ECTR than those in patients who underwent OCTR (OR, 4.87; 95% CI, 1.37 to 17.25; $P = 0.01$; $n = 2320$; fixed-effects model, with a heterogeneity of $I^2 = 0\%; P = 0.98$) (Fig. 13); however, the studies provided evidence that the presence of permanent nerve injury was not significantly different between the two groups (OR, 1.93; 95% CI, 0.58 to 6.40; $P = 0.28$; $n = 2320$; fixed-effects model, with a heterogeneity of $I^2 = 29\%; P = 0.24$) (Fig. 14). The rates of scar-related complications (scar hypertrophy, scar hyperesthesia, scar pain) were lower in patients who underwent ECTR than those in patients who underwent OCTR (OR, 0.20; 95% CI, 0.07 to 0.59; $P = 0.004$; $n = 2320$; fixed-effects model with a heterogeneity of $I^2 = 0\%; P = 0.90$) (Fig. 15). Other complications, such as hematoma, wound infection, superficial palmar arch injury, persistent symptoms, pillar pain, reflex sympathetic dystrophy, and tendon injury, were not significantly different between the two groups. The summary of all outcome variables is shown in Table 2.

3.4. Publication bias

The funnel plot method and Egger’s test were used to evaluate publication bias in the included literature. The funnel plot shape and Egger’s test ($P = 0.869$) did not reveal any obvious asymmetry
(Fig. 16), indicating no overt publication bias in the analysis of complications.

4. Discussion

Since the development of ECTR by Chow (12) and Okutsu et al. [13] in 1989, there has been controversy regarding the superiority of ECTR over OCTR. Accordingly, many original articles have been published on this issue; moreover, several meta-analyses have compared ECTR with OCTR as treatment options for CTS [14,15,21-23,51-53]. However, previous meta-analyses included fewer studies than ours, did not classify the data into subgroups according to different follow-up times, featured only a few assessments of patient outcome, and included central tendency data but not standard deviation. Therefore, we performed a large sample-size meta-analysis of published results to compare the effectiveness and safety of the two surgical approaches. The publication bias in this meta-analysis was also minimal, as demonstrated by the results of the funnel plot analysis and Egger’s test.

Our meta-analysis reviewed 28 RCTs that consisted of 2,320 idiopathic CTS hands treated with OCTR or ECTR. The results clearly indicated that there were no significant between-group differences in the operative time, grip strength, BCTQ-S score, BCTQ-F score, digital sensation scores, and the presence of permanent nerve injury. However, the ECTR group exhibited several clinically important advantages over the OCTR group, including higher patient satisfaction rates, greater key pinch strengths, earlier RTW times, and fewer scar-related complications.

Consistent with the present results, previous studies demonstrated that the satisfaction rates of patients in the ECTR group were higher than those of patients in the OCTR group [24,47,54]. Compared with the standard open approach, reduction in scar tenderness after endoscopic release is plausible, owing to the generally small incisions, reduced scarring, mild wound-related complications [34], and improvements in the major functional outcomes (key pinch strength, activities of daily living, and RTW) [32]. However, it should be noted that when assessing the patient satisfaction rates, a portion of the data published by Zhang et al. [34] exhibited high heterogeneity. Therefore, these data were excluded from the present meta-analysis. The high heterogeneity was mainly because of the fact that the data compared mini-incisions with endoscopic incisions. Mini-incisions are not
directly comparable to the standard incisions in OCTR, as they yield a better appearance and tend to have fewer wound-related complications than standard incisions [55].

Herein, the key pinch strength of patients in the ECTR group was significantly greater than that in the OCTR group at 3 months postoperatively [9,32]. Additionally, previous studies reported that OCTR was associated with considerable morbidity, including increased and prolonged scar tenderness [11]. Furthermore, other studies revealed that patients who underwent ECTR experienced fewer limitations in their ability to perform daily life activities than did patients who underwent an open technique [36,56-58]. Michelotti et al. [54] reported early differences in grip and pinch strength after ECTR; however, data were lost as the follow-up duration increased. Further studies should include a more uniform follow-up duration, and additional controlled studies with longer follow-up durations are required to clarify the effects of each technique on activities of daily living.

The finding of our meta-analysis of RCTs suggest that patients treated with ECTR returned to work or daily activities earlier than those treated with OCTR. Consistent with our results, Vasiliadis et al. [22] and Paryavi et al. [52] reported that patients who underwent ECTR experienced less surgical trauma than those who underwent an open technique, and this resulted in less time off work, faster recovery, and better performance of daily activities. However, regarding the RTW data, we noticed that divergences between the studies resulted in large between-study heterogeneity. A possible explanation for this large variability is that the flexibility required for the work and the nature of the work and daily activities may have been substantially different in the included studies. Furthermore, while Sanati et al. [53] demonstrated the superiority of minimally invasive techniques over conventional open release in terms of recovery time, they highlighted the remarkable inconsistencies in how RTW as an outcome measure was examined across studies. Nevertheless, the effects of such inconsistencies were rather small when only RCTs were considered, similar to that observed in our study. Compared with open release, patients undergoing endoscopic release can return to work and their daily activities sooner.

Our meta-analysis revealed that lower scar-related complication rates and better healing were achieved in the ECTR group than in the OCTR group. This may be because of the extended incision in...
the palm made during OCTR that may extend the immobilization time and increase postoperative pain and the risk for hypertrophic or hypersensitive scars [22]. In contrast, ECTR uses a small incision and divides the transverse carpal ligament from below, thereby preserving the overlying skin and muscle and resulting in fewer minor complications [59,60], particularly those related to cutaneous scars. However, previous studies demonstrated that ECTR is associated with more nerve injury; therefore, the technique is less favorable owing to its higher risk of iatrogenic injury to the cutaneous branch of the median nerve [15,19,34]. Contrary to expectations, our study did not find a significant difference in the occurrence of permanent nerve injury between the two surgical approaches; furthermore, most noted nerve injuries were transient, and patients still achieved full recovery after surgery [37,38,50,61]. Moreover, Martin et al. [62] developed a novel endoscopic system which may avoid the transient nerve injury occurring with other ECTR methods.

4.1. Limitations

This study had two limitations. First, subgroup analyses of the various ECTR techniques (one-portal and two-portal techniques) and OCTR techniques (mini-incision and long incision) were not performed. Although the use of different techniques may be associated with different outcomes, we were unable to perform subgroup analyses because of the rather small number of studies and limited available data. Second, although we included only RCTs, all the trials had methodological flaws, including unblinded assessments of outcomes.

Nevertheless, our study is novel since it includes the largest number of RCTs to compare ECTR and OCTR techniques. Furthermore, this is the first study to group results into different follow-up times and assess different patient outcomes, thus making the data more comparable. This study is also the first to demonstrate that ECTR is associated with better patient outcomes; we found that after careful manipulation during endoscopic surgery, ECTR can substitute OCTR.

5. Conclusions

The present meta-analysis determined that ECTR was superior to OCTR in terms of higher satisfaction rates, improved key pinch strengths, earlier RTW times, and fewer scar-related complications. Our findings suggest that patients with CTS can be effectively managed with ECTR; however, the
possibility of transient nerve injury should be considered.

List Of Abbreviations
CTS: carpal tunnel syndrome
OCTR: open carpal tunnel release
ECTR: endoscopic carpal tunnel release
RCT: randomized controlled trial
PRISMA: Preferred Reporting Items for Systematic Reviews and Meta-Analyses
BCTQ-S: Boston Carpal Tunnel Questionnaire Symptom Severity Scale
BCTQ-F: Boston Carpal Tunnel Questionnaire Functional Status Scale
RTW: return to work
GRADE: Grading of Recommendations Assessment, Development, and Evaluation
MD: mean difference
CI: confidence interval
OR: odds ratio
SE: standard error

Declarations
Ethics approval and consent to participate
This article does not contain any studies involving human participants conducted by any of the authors.

Consent for publication
Not applicable.

Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Competing interests
The authors declare that they have no competing interests.

Funding
This work was supported by the Jilin Scientific and Technological Development Program [grant number 20190905003SF]. The funding institution was responsible for the provision of literature database resources and literature purchase. The funding body had no role in the design of the study, the collection, analysis, and interpretation of data, or the writing of the manuscript.

Acknowledgments
Not applicable.

Author information
Yueying Li is the first author

Affiliations
Department of Hand Surgery, China-Japan Union Hospital of Jilin University, No. 126 Xiantai Street, Changchun, Jilin 130033, P.R. China
Yueying Li, Guangzhi Wu, Shusen Cui, Zhan Zhang & Xiaosong Gu
Department of Orthopedics, China-Japan Union Hospital of Jilin University, No. 126 Xiantai Street, Changchun, Jilin 130033, P.R. China
Wenqi Luo

Author’s Contributions
YL analyzed and interpreted the data and was a major contributor in writing the manuscript. LW collected the data. GW analyzed and interpreted the data. SC revised the article critically. XG designed the study and supervised the study. ZZ designed the study and revised the article critically. All authors read and approved the final manuscript.

Corresponding authors
Correspondence to Xiaosong Gu or Zhan Zhang

References
(1) Devana SK, Jensen AR, Yamaguchi KT, D'Oro A, Buser Z, Wang JC, et al. Trends and complications in open versus endoscopic carpal tunnel release in private payer and medicare patient populations. Hand (N Y). 2019;14:455-61. doi:10.1177/1558944717751196.
(2) Atroshi I, Gummesson C, Johnsson R, Ornstein E, Ranstam J, Rosén I. Prevalence of carpal
tunnel syndrome in a general population. JAMA. 1999;282:153–8. doi:10.1001/jama.282.2.153.

(3) Bongers FJ, Schellevis FG, van den Bosch WJ, van der Zee J. Carpal tunnel syndrome in general practice (1987 and 2001): incidence and the role of occupational and non-occupational factors. Br J Gen Pract. 2007;57:36–9.

(4) Burton CL, Chen Y, Chesterton LS, Van der Windt DA. Trends in the prevalence, incidence, and surgical management of carpal tunnel syndrome between 1993 and 2013: an observational analysis of UK primary care records. BMJ Open. 2018;8:e020166. doi:10.1136/bmjopen-2017-020166.

(5) Middleton SD, Anakwe RE. Carpal tunnel syndrome. BMJ. 2014;349:g6437. doi:10.1136/bmj.g6437.

(6) Gerritsen AA, de Vet HC, Scholten RJ, Bertelsmann FW, de Krom MC, Bouter LM. Splinting vs. surgery in the treatment of carpal tunnel syndrome: a randomized controlled trial. JAMA. 2002;288:1245–51. doi:10.1001/jama.288.10.1245.

(7) Chesterton LS, Blagojevic-Bucknall M, Burton C, Dziedzic KS, Davenport G, Jowett SM, et al. The clinical and cost-effectiveness of corticosteroid injection versus night splints for carpal tunnel syndrome (INSTINCTS trial): an open-label, parallel group, randomised controlled trial. Lancet. 2018;392:1423–1433. doi:10.1016/S0140-6736(18)31572-1.

(8) Ebenbichler GR1, Resch KL, Nicolakis P, Wiesinger GF, Uhl F, Ghanem AH, et al. Ultrasound treatment for treating the carpal tunnel syndrome: randomised "sham" controlled trial. BMJ. 1998;316:731–5. doi:10.1136/bmj.316.7133.731.

(9) Atroshi I, Larsson GU, Ornstein E, Hofer M, Johnsson R, Ranstam J. Outcomes of endoscopic surgery compared with open surgery for carpal tunnel syndrome among employed patients: randomised controlled trial. BMJ. 2006;332:1473. doi:10.1136/bmj.38863.632789.1F.

(10) Phalen GS. The carpal-tunnel syndrome, Seventeen years' experience in diagnosis and treatment of six hundred fifty-four hands. J Bone Joint Surg Am. 1966;48:211–28.

(11) Boya H, Özcan Ö, Özteki NHH. Long-term complications of open carpal tunnel release. Muscle Nerve. 2008;38:1443–6. doi:10.1002/mus.21068.

(12) Chow JC. Endoscopic release of the carpal ligament: a new technique for carpal tunnel
syndrome. Arthroscopy, 1989;5:19–24. doi:10.1016/0749-8063(89)90085-6.

(13) Okutsu I, Ninomiya S, Takatori Y, Ugawa Y. Endoscopic management of carpal tunnel syndrome. Arthroscopy. 1989;5:11–8. doi:10.1016/0749-8063(89)90084-4.

(14) Chen L, Duan X, Huang X, Lv J, Peng K, Xiang Z. Effectiveness and safety of endoscopic versus open carpal tunnel decompression. Arch Orthop. Trauma Surg. 2014;134:585–93. doi:10.1007/s00402-013-1898-z.

(15) Sayegh ET, Strauch RJ. Open versus endoscopic carpal tunnel release: a meta-analysis of randomized controlled trials. Clin Orthop Relat Res. 2015;473:1120–32. doi:10.1007/s11999-014-3835-z.

(16) Cobb TK, Knudson GA, Cooney WP. The use of topographical landmarks to improve the outcome of Agee endoscopic carpal tunnel release. Arthroscopy. 1995;11:165–172. doi:10.1016/0749-8063(95)90062-4.

(17) Concannon MJ, Brownfield ML, Puckett CL. The incidence of recurrence after endoscopic carpal tunnel release. Plast Reconstr Surg. 2000;105:1662–5. doi:10.1097/00006534-200004050-00010.

(18) Makowiec RL, Nagle DJ, Chow JC. Outcome of first-time endoscopic carpal tunnel release in a teaching environment. Arthroscopy. 2002;18:27–31. doi:10.1053/jars.2002.29903.

(19) Murphy Jr. RX, Jennings JF, Wukich DK. Major neurovascular complications of endoscopic carpal tunnel release. J Hand Surg Am. 1994;19:114–8. doi:10.1016/0363-5023(94)90233-X.

(20) Chung KC, Walters MR, Greenfield ML, Chernew M. Endoscopic versus open carpal tunnel release: a cost-effectiveness analysis. Plast Reconstr Surg. 1998;102:1089–99. doi:10.1097/00006534-199809040-00026.

(21) Zuo D, Zhou Z, Wang H, Liao Y, Zheng L, Hua Y, et al. Endoscopic versus open carpal tunnel release for idiopathic carpal tunnel syndrome: a meta-analysis of randomized controlled trials. J Orthop Surg Res. 2015;10:12. doi:10.1186/s13018-014-0148-6.

(22) Vasiliadis HS, Nikolakopoulou A, Shrier I, Lunn MP, Brassington R, Scholten RJ, et al. Endoscopic and open release similarly safe for the treatment of carpal tunnel syndrome. A systematic review and meta-analysis. PloS One. 2015;10:e0143683. doi:10.1371/journal.pone.0143683.
(23) Hu K, Zhang T, Xu W. Intraindividual comparison between open and endoscopic release in bilateral carpal tunnel syndrome: a meta-analysis of randomized controlled trials. Brain Behav. 2016;6:e00439. doi:10.1002/brb3.439.

(24) Martínez-Catasús A, Lobo-Escolar L, García-Bonet J, Corrales-Rodríguez M, Pasarín-Martínez A, Berlanga-de-Mingo D. Comparison between single portal endoscopic, 1-cm open carpal tunnel release. Hand Surg Rehabil. 2019;38:202–6. doi:10.1016/j.hansur.2019.02.003.

(25) Atroshi I, Hofer M, Larsson GU, Ranstam J, Extended follow-up of a randomized clinical trial of open vs. endoscopic release surgery for carpal tunnel syndrome. JAMA. 2015;314:1399–401. doi:10.1001/jama.2015.12208.

(26) Moher D, Liberati A, Tetzlaff J, Altman DG, PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. Ann Intern Med. 2009;151:264–9. doi:10.7326/0003-4819-151-4-200908180-00135.

(27) Guyatt GH, Oxman AD, Schünemann Hj, Tugwell P, Knottnerus A. GRADE guidelines: a new series of articles in the Journal of Clinical Epidemiology. J Clin Epidemiol. 2011;64:380–2. doi:10.1016/j.jclinepi.2010.09.011.

(28) Ferdinand RD, MsacLean JG. Endoscopic versus open carpal tunnel release in bilateral carpal tunnel syndrome. A prospective, randomised, blinded assessment. J Bone Joint Surg Br. 2002;84:375–9. doi:10.1302/0301-620X.84B3.12224.

(29) Kang HJ, Koh IH, Lee TJ, Choi YR. Endoscopic carpal tunnel release is preferred over mini-open despite similar outcome: a randomized trial. Clin Orthop Relat Res. 2013;471:1548–54. doi:10.1007/s11999-012-2666-z.

(30) Wong KC, Hung LK, Ho PC, Wong JM. Carpal tunnel release. A prospective, randomised study of endoscopic versus limited-open methods. J Bone Joint Surg Br. 2003;85:863–8. doi:10.1302/0301-620X.85B6.13759.

(31) Zhao H, Zhao Y, Tian Y, Yang B, Qiu GX. Comparison of endoscopic versus open surgical treatment of carpal tunnel syndrome. Zhongguo Yi Xue Ke Xue Yuan Xue Bao 2004;26:657–60.

(32) Brown RA, Gelberman RH, Seiler 3rd JG, Abrahamsson SO, Weiland AJ, Urbaniak JR, et al.
Carpal tunnel release. A prospective, randomized assessment of open and endoscopic methods. J Bone Joint Surg Am. 1993;75:1265–75. doi:10.2106/00004623-199309000-00002.

(33) Trumble TE, Diao E, Abrams RA, Gilbert-Anderson MM. Single-portal endoscopic carpal tunnel release compared with open release: a prospective, randomized trial. J Bone Joint Surg Am. 2002;84:1107–15. doi:0.2106/00004623-200207000-00003.

(34) Zhang X, Huang X, Wang X, Wen S, Sun J, Shao X. A randomized comparison of double small, standard, and endoscopic approaches for carpal tunnel release. Plast Reconstr Surg. 2016;138:641–7. doi:10.1097/PRS.0000000000002511.

(35) Tian Y, Zhao H, Wang T. Prospective comparison of endoscopic and open surgical methods for carpal tunnel syndrome, Chin Med Sci J. 2007;22:104–7.

(36) Aslani HR, Alizadeh K, Eajazi A, Karimi A, Karimi MH, Zaferani Z, et al. Comparison of carpal tunnel release with three different techniques. Clin Neurol Neurosurg. 2012;114:965–8. doi:10.1016/j.clineuro.2012.02.017.

(37) Jacobsen MB, Rahme H. A prospective, randomized study with an independent observer comparing open carpal tunnel release with endoscopic carpal tunnel release. J Hand Surg Br. 1996;21:202–4. doi:10.1016/s0266-7681(96)80097-0.

(38) Saw NL, Jones S, Shepstone L, Meyer M, Chapman PG, Logan AM. Early outcome and cost-effectiveness of endoscopic versus open carpal tunnel release: a randomized prospective trial. J Hand Surg Br. 2003;28:444–9. doi:10.1016/s0266-7681(03)00097-4.

(39) Agee JM, McCarroll HR Jr, Tortosa RD, Berry DA, Szabo RM, Peimer CA. Endoscopic release of the carpal tunnel: a randomized prospective multicenter study. J Hand Surg Am. 1992;17:987–95. doi:10.1016/s0363-5023(09)91044-9.

(40) Dumontier C, Sokolow C, Leclerq C, Chauvin P. Early results of conventional versus two-portal endoscopic carpal tunnel release. A prospective study. J Hand Surg Br. 1995;20:658–62. doi:10.1016/s0266-7681(05)80130-5.

(41) Ejiri S, Kikuchi S, Maruya M, Sekiguchi Y, Kawakami R, Konno S. Short-term results of endoscopic (Okutsu method) versus palmar incision open carpal tunnel release: a prospective
randomized controlled trial. Fukushima J Med Sci. 2012;58:49-59. doi:10.5387/fms.58.49.

(42) Erdmann MW. Endoscopic carpal tunnel decompression. J Hand Surg Br. 1994;19:5-13. doi:10.1016/0266-7681(94)90038-8.

(43) Gümüştaş SA, Ekmekçi B, Tosun HB, Orak MM, Bekler Hİ. Similar effectiveness of the open versus endoscopic technique for carpal tunnel syndrome: a prospective randomized trial. Eur J Orthop Surg Traumatol. 2015;25:1253-60. doi:10.1007/s00590-015-1696-0.

(44) Larsen MB, Sørensen Al, Crone KL, Weis T, Boeckstyns ME. Carpal tunnel release: a randomized comparison of three surgical methods. J Hand Surg Eur Vol. 2013;38:646-50. doi:10.1177/1753193412475247.

(45) Macdermid JC, Richards RS, Roth JH, Ross DC, King GJ. Endoscopic versus open carpal tunnel release: a randomized trial. J Hand Surg Am. 2003;28:475-80. doi:10.1053/jhsu.2003.50080.

(46) Mackenzie Dj, Hainer R, Wheatley Mj. Early recovery after endoscopic vs. short-incision open carpal tunnel release. Ann Plast Surg. 2000;44:601-4. doi:10.1097/00000637-200044060-00004.

(47) Michelotti BM, Vakharia KT, Romanowsky D, Hauck RM. A prospective, randomized trial comparing open and endoscopic carpal tunnel release within the same patient. Hand (N Y). 2018;21:1558944718812129. doi:10.1177/1558944718812129.

(48) Oh WT, Kang HJ, Koh IH, Jang JY, Choi YR. Morphologic change of nerve and symptom relief are similar after mini-incision and endoscopic carpal tunnel release: a randomized trial. BMC Musculoskelet Disord. 2017;18:65. doi:10.1186/s12891-017-1438-z.

(49) Rab M, Grünbeck M, Beck H, Haslik W, Schrögendorfer KF, Schiefer HP, et al. Intra-individual comparison between open and 2-portal endoscopic release in clinically matched bilateral carpal syndrome. J Plast Reconstr Aesthet Surg. 2006;59:730–6. doi:10.1016/j.bjps.2005.11.018.

(50) Sennwald GR, Benedetti R. The value of one-portal endoscopic carpal tunnel release: a prospective randomized study. Knee Surg Sports Traumatol Arthrosc. 1995;3:113-116. doi:10.1007/BF01552386.

(51) Thoma A, Veltri K, Haines T, Duku E. A meta-analysis of randomized controlled trials comparing endoscopic and open carpal tunnel decompression. Plast Reconstr Surg. 2004;114:1137-
46. https://doi:10.1097/01.prs.0000135850.37523.d0.

(52) Paryavi E, Zimmerman RM, Means Jr. KR. Endoscopic compared with open operative treatment of carpal tunnel syndrome. JBJS Rev. 2016;4. doi:10.2106/JBJS.RVW.15.00071.

(53) Sanati KA, Mansouri M, Macdonald D, Ghafghazi S, Macdonald E, Yadegarfar G. Surgical techniques and return to work following carpal tunnel release: a systematic review and meta-analysis. J Occup Rehabil. 2011;21:474–81. doi:10.1007/s10926-011-9310-8.

(54) Michelotti B, Romanowsky D, Hauck RM. Prospective, randomized evaluation of endoscopic versus open carpal tunnel release in bilateral carpal tunnel syndrome: an interim analysis. Ann Plast Surg. 2014;73 Suppl 2:S157–S160. doi:10.1097/SAP.0000000000000203.

(55) Li G, Kong L, Kou N, Wang Y, Yu K, Bai J, et al. Tian, The comparison of limited-incision versus standard-incision in treatment of carpal tunnel syndrome: a meta-analysis of randomized controlled trials. Medicine (Baltimore). 2019;98:e15372. doi:10.1097/MD.00000000000015372.

(56) Gellman H, Kan D, Gee V, Kuschner SH, Botte MJ. Analysis of pinch and grip strength after carpal tunnel release. J Hand Surg Am. 1989;14:863–4. doi:10.1016/s0363-5023(89)80091-7.

(57) Cotton P. Symptoms may return after carpal tunnel surgery. JAMA. 1991;265:1922–5. doi:10.1001/jama.265.15.1922b.

(58) Macfarlane D, Williams TG. Efficacy of provocative tests for diagnosis of carpal tunnel syndrome. Lancet. 1990;335:727. doi:10.1016/0140-6736(90)90843-t.

(59) Kim PT, Lee HJ, Kim TG, Jeon IH. Jeon, Current approaches for carpal tunnel syndrome. Clin Orthop Surg. 2014;6:253–7. doi:10.4055/cios.2014.6.3.253.

(60) Padua L, Coraci D, Erra C, Pazzaglia C, Paolasso I, Loreti C, et al. Carpal tunnel syndrome: clinical features, diagnosis, and management. Lancet Neurol. 2016;15:1273–84. doi:10.1016/S1474-4422(16)30231-9.

(61) Atroshi I, Hofer M, Larsson GU, Ornstein E, Johnsson R, Ranstam J. Open compared with 2-portal endoscopic carpal tunnel release: a 5-year follow-up of a randomized controlled trial. J Hand Surg Am. 2009;34:266–72. doi:10.1016/j.jhسا.2008.10.026.

(62) Martin KD, Dützmann S, Sobottka SB, Rambow S, Mellerowicz HA, Pinzer T, et al. Retractor-
endoscopic nerve decompression in carpal and cubital tunnel syndromes: outcomes in a small series.

World Neurosurg. 2014;82:e361–e370. doi:10.1016/j.wneu.2013.09.026.

(63) Taleisnik J. The palmar cutaneous branch of the median nerve and the approach to the carpal tunnel. J Bone Joint Surg Am. 1973;55:1212-1217.

(64) Menon J. Endoscopic carpal tunnel release: a single-portal technique. Contemp Orthop. 1993;26:109-16.

Tables

Table 1. Study characteristics of the randomized controlled trials included in the meta-analysis

| Author            | Year | Region | Groups, sample sizes, and techniques                                      | Follow-up  |
|-------------------|------|--------|---------------------------------------------------------------------------|------------|
| Agee et al. [39]  | 1992 | US     | OCTR (n=65): conventional release                                          | 1, 2, 3, 6,|
|                   |      |        | ECTR (n=82): one-portal (2 cm)                                             |            |
| Atroshi et al. [9]| 2006 | Sweden | OCTR (n=65): 4 cm                                                          | 3, 6 week  |
|                   |      |        | ECTR (n=63): two-portal (1 cm)                                             | 3, 12 mor  |
| Atroshi et al. [61]| 2009 | Sweden | OCTR (n=63): 4 cm                                                          | 1, 5 year  |
|                   |      |        | ECTR (n=63): two-portal (1 cm)                                             |            |
| Atroshi et al. [25]| 2015 | Sweden | OCTR (n=61): 4 cm                                                          | 1, 11-16   |
|                   |      |        | ECTR (n=63): two-portal (1 cm)                                             |            |
| Aslani et al. [36]| 2012 | Iran   | OCTR (n=36): conventional release                                          | 4 weeks;   |
|                   |      |        | OCTR (n=28): mini-incision                                                  | 4 months   |
|                   |      |        | ECTR (n=32): two-portal (length NR)                                        |            |
| Brown et al. [32] | 1993 | US     | OCTR (n=82): conventional release                                          | 3, 6, 12 w|
|                   |      |        | ECTR (n=78): two-portal (2 cm)                                             |            |
| Dumontier et al. [40]| 1995 | France | OCTR (n=40): conventional release                                          | 2 weeks;   |
|                   |      |        | ECTR (n=56): two-portal (Chow [12])                                        | 1, 3, 6 mc |
| Ejiri et al. [41] | 2012 | Japan  | OCTR (n=50): 3 cm vertical incision                                         | 1, 3 mont  |
|                   |      |        | ECTR (n=50): one-portal (Okutsu et al. [13])                              |            |
| Authors                  | Year | Location | OCTR n=52: (short length) | ECTR n=53: (Chow [12]) | OCTR n=25: | ECTR n=25: |
|--------------------------|------|----------|--------------------------|------------------------|-----------|-----------|
| Erdmann                  | 1994 | UK       | NR                       | two-portal             | NR        | one-portal|
| Ferdinand and MacLean    | 2002 | UK       | NR                       | two-portal (Chow [12]) | 6, 12, 26 | 13 month  |
| Gümüştaş et al.          | 2015 | Turkey   | NR                       | two-portal (Chow [12]) | 6 months  |
| Jacobson and Rahme       | 1996 | Sweden   | conventional release     | two-portal (Chow [12]) | 2, 6, 24 w|
| Kang et al.              | 2013 | South Korea | mini-incision (1.5 cm) | one-portal (Agee et al. [39]) | 3 months  |
| Larsen et al.            | 2013 | Denmark  | classic incision, 7 cm   | one-portal (Menon [64]) | 1, 2, 3, 6, 12 months |
| Macdermid et al.         | 2003 | Canada   | two-portal (Chow [12])   | 1, 6, 12 w             |
| Mackenzie et al.         | 2000 | US       | 2.5 cm palmar incision   | one-portal (Agee et al. [39]) | 1, 2, 4 week |
| Martínez et al.          | 2019 | Spain    | 1 cm mini-incision       | one-portal (Menon [64]) | 1 week; 1, 6, 12 months |
| Michelotti et al.        | 2014 | US       | 3 cm palmar incision     | one-portal (Agee et al. [39], 1.5 cm) | 2, 4, 8, 12 |
| Michelotti et al.        | 2018 | US       | 3 cm palmar incision     | one-portal (Agee et al. [39], 1.5 to 2 cm) | 2, 4, 8, 12 |
| Oh et al.                | 2017 | South Korea | mini-incision (1.5 cm) | one-portal (Agee et al. [39], 1.5 cm) | 24 weeks |
| Rab et al.               | 2006 | Austria  | two mini-incisions       | two-portal (Chow [12]) | 2, 4, 6, 12 months |
| Saw et al.               | 2003 | UK       | 2 cm palmar incision     | one-portal (Agee et al. [39]) | 1, 3, 6, 12 |
| Sennwald and             | 1995 | Switzerland | Sennwald incision      | 4, 8, 12 w             |
| Study                  | Year | Location | OCTR (n=) | Procedure | ECTR (n=) | Procedure | Follow-up |
|------------------------|------|----------|-----------|------------|-----------|-----------|-----------|
| Benedetti [50]         |      |          |           |            |           |           | 2 years   |
| Tian et al. [35]       | 2007 | China    | 36        | S-shaped incision | 34        | one-portal (Okutsu et al. [13], 1 cm) | 2 years   |
| Trumble et al. [33]    | 2002 | US       | 95        | palmar incision (3~4 cm) | 97        | one-portal (Agee et al. [39], 1 cm) | 2, 4, 8, 12 years |
| Wong et al. [30]       | 2003 | Hong Kong| 30        | mini-incision | 30        | two-portal (Chow [12]) | 2, 4, 8, 12 months |
| Zhang et al. [34]      | 2016 | China    | 72        | double small incision | 65        | standard incision (5~7 cm) | 3 years |
| Zhao et al. [31]       | 2004 | China    | 21        | S-shaped incision | 26        | one-portal (Okutsu et al. [13], 1 cm) | 2 years |

2PD: Two-point Discrimination; ADL: activities of daily living; APB: abductor pollicis brevis; BCTQ-F: Boston Carpal Tunnel Questionnaire Functional Status Scale; BCTQ-S: Boston Carpal Tunnel Questionnaire Symptom Severity Scale; CMAP: compound muscle action potential; CSA: cross-sectional area; CSA-I: inlet at the distal wrist crease level; CSA-M: the middle of the tunnel at the level of the pisiform; CSA-O: the tunnel outlet at the level of the hamate hook; DASH: Disability of Arm, Shoulder, and Hand Questionnaire; DL: distal latency; ECTR: endoscopic carpal tunnel release; EMG: electromyography; NCV: nerve conduction velocity; NR: not reported; OCTR: open carpal tunnel release; RTW: return to work; SCV: sensory conduction velocity; SF-12: 12-Item Short Form Health Survey; SF-36: 36-Item Short Form Health Survey; SW: Semmes-Weinstein; UK: United Kingdom; US: United States; VAS: Visual Analog Scale.

Table 2. Summary estimates of outcome variables in the current study
| Outcome                          | No. of studies | Heterogeneity $I^2$ (%) | Pooled odds ratio† | Mean difference |
|---------------------------------|----------------|------------------------|-------------------|-----------------|
| Operative time                  | 4              | 99                     | NA                | -5.81           |
| Grip strength                   | 2              | 0                      | NA                | 1.99            |
| BCTQ-S score                    | 3              | 92                     | NA                | 0.15            |
| BCTQ-F score                    | 3              | 91                     | NA                | 0.17            |
| SW monofilament test score      | 2              | 0                      | NA                | 0.06            |
| 2PD test score                  | 3              | 35                     | NA                | -0.16           |
| Satisfaction rate               | 2              | 0                      | NA                | 3.13            |
| Key pinch strength              | 2              | 0                      | NA                | 0.79            |
| Return to work                  | 4              | 98                     | NA                | -7.25           |
| Complications                   | 25             | 16                     | 1.06              | NA              |
| Transient nerve injury          | 25             | 0                      | 4.87              | NA              |
| Permanent nerve injury          | 25             | 29                     | 1.93              | NA              |
| Scar-related complications      | 25             | 0                      | 0.2               | NA              |
| Hematoma                        | 25             | 0                      | 1.60              | NA              |
| Wound infection                 | 25             | 0                      | 0.53              | NA              |
| Superficial palmar arch injury  | 25             | NA                     | 3.07              | NA              |
| Persistent symptoms             | 25             | 0                      | 2.17              | NA              |
| Pillar pain                     | 25             | 35                     | 0.95              | NA              |
| Reflex sympathetic dystrophy    | 25             | 0                      | 0.40              | NA              |
| Tendon injury                   | 25             | NA                     | 0.26              | NA              |

2PD: Two-point Discrimination; BCTQ-F: Boston Carpal Tunnel Questionnaire Functional Status Scale; BCTQ-S: Boston Carpal Tunnel Questionnaire Symptom Severity Scale; ECTR: endoscopic carpal tunnel release; NA: not applicable; OCTR: open carpal tunnel release; SW: Semmes-Weinstein.

*Heterogeneity test: $I^2 > 50\%$, random-effects analysis model; $I^2 < 50\%$, fixed-effects analysis model.

†If odds ratio $>1$, favors ECTR; if odds ratio $<1$, favors OCTR.

‡If mean difference $>0$, favors ECTR; if mean difference $<0$, favors OCTR.

Figures
References identified in database search (n=5654)
PubMed (n=1416), EMBASE (n=1755), Cochrane Library (n=248), Web of Science (n=1130), Medline (n=1105), Additional records identified through reference lists (n=0)

Records after duplicates removed (n=2248)

Screening of titles and abstracts (n=2248)
Records excluded (n=2145)

Full-text articles assessed for eligibility (n=103)

Full-text articles excluded, with reasons (n=75)
Nonrandomized study design (n=32)
Review, abstract, letter, congress, case report (n=14)
No full text available (n=6)
Other languages (n=5)
Limited information (n=18)

Studies included in quantitative synthesis (meta-analysis) (n=28)
27 articles in English, 1 article in Chinese

Figure 1
Flow diagram of the study selection process.
| Study             | Random sequence generation | Allocation concealment (select) | Blinding of participants and personnel | Blinding of outcome assessment | Incomplete outcome data (attrition) | Selective reporting (reporting bias) | Other bias |
|-------------------|-----------------------------|---------------------------------|----------------------------------------|-------------------------------|-----------------------------------|-------------------------------------|------------|
| Agee 1992         | ?                           | ?                               | -                                      | -                             | +                                 | +                                   | +          |
| Aslani 2012       | ?                           | ?                               | -                                      | -                             | +                                 | +                                   | +          |
| Atroshi 2006      | +                           | ?                               | -                                      | -                             | +                                 | +                                   | +          |
| Atroshi 2009      | +                           | ?                               | -                                      | -                             | +                                 | +                                   | +          |
| Atroshi 2015      | +                           | ?                               | -                                      | -                             | +                                 | +                                   | +          |
| Brown 1993        | +                           | ?                               | -                                      | -                             | -                                 | +                                   | +          |
| Dumontier 1995    | -                           | ?                               | -                                      | -                             | -                                 | +                                   | +          |
| Ejiri 2012        | +                           | -                               | -                                      | -                             | +                                 | +                                   | +          |
| Erdmann 1994      | +                           | ?                               | ?                                      | ?                             | ?                                 | +                                   | +          |
| Ferdinand 2002    | +                           | -                               | +                                      | +                             | +                                 | +                                   | +          |
| Gümüştaş 2015     | +                           | ?                               | -                                      | +                             | +                                 | +                                   | +          |
| Jacobsen 1996     | -                           | -                               | -                                      | +                             | +                                 | +                                   | +          |
| Study              | A | B | C | D | E | F | G | H |
|--------------------|---|---|---|---|---|---|---|---|
| Kang 2013          | ? |   |   |   |   |   |   |   |
| Larsen 2013        |   |   |   |   |   |   |   |   |
| Macdermid 2003     |   |   |   |   |   |   |   |   |
| Mackenzie 2000     |   |   |   |   |   |   |   |   |
| Martínez-Catasús 2019 | | | | | | | | |
| Michelotti 2014    |   |   |   |   |   |   |   |   |
| Michelotti 2018    |   |   |   |   |   |   |   |   |
| Oh 2017            |   |   |   |   |   |   |   |   |
| Rab 2006           |   |   |   |   |   |   |   |   |
| Saw 2003           |   |   |   |   |   |   |   |   |
| Sennwald 1995      |   |   |   |   |   |   |   |   |
| Tian 2007          |   |   |   |   |   |   |   |   |
| Trumble 2002       |   |   |   |   |   |   |   |   |
| Wong 2003          |   |   |   |   |   |   |   |   |
| Zhang 2016         |   |   |   |   |   |   |   |   |
| zhao 2004          |   |   |   |   |   |   |   |   |

Figure 2

Risk of bias assessment for included randomized controlled trials.
Figure 3

Comparison of operative time between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.

| Study or Subgroup | ECTR Mean | SD | Total | OCTR Mean | SD | Total | Weight | Mean Difference | IV, Random, 95% CI |
|-------------------|-----------|----|-------|-----------|----|-------|--------|-----------------|-------------------|
| Zhao 2004         | 13        | 2  | 26    | 40        | 5  | 21    | 25.0%  | -27.00 [-29.27, -24.73] |
| Wong 2003         | 12.9      | 4.9| 30    | 12.9      | 5.1| 30    | 24.9%  | 0.00 [2.53, 2.53]   |
| Kang 2013         | 7.5       | 3.68| 52    | 6.8       | 2.58| 52    | 25.1%  | 0.70 [-0.52, 1.92]  |
| Ferdinand 2002    | 13        | 4  | 25    | 10        | 2  | 25    | 25.0%  | 3.00 [1.25, 4.75]   |
| Total (95% CI)    | 133       |    |       | 128       |    |       | 100.0% | -5.81 [-17.85, 6.23] |

Heterogeneity: Tau^2 = 149.98; Chi^2 = 513.93, df = 3 (P < 0.00001); P = 99%
Test for overall effect: Z = 0.95 (P = 0.34)

Figure 4

Comparison of grip strength between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.

| Study or Subgroup | ECTR Mean | SD | Total | OCTR Mean | SD | Total | Weight | Mean Difference | IV, Fixed, 95% CI |
|-------------------|-----------|----|-------|-----------|----|-------|--------|-----------------|-------------------|
| 7.1.1 3m          |           |    |       |           |    |       |        |                 |                   |
| Atroshi 2006      | 31.5      | 11 | 63    | 29.9      | 11 | 65    | 40.6%  | 1.60 [-2.21, 5.41] |
| Brown 1993        | 23.13     | 10.84| 84    | 20.87     | 10.02| 85    | 59.4%  | 2.26 [-0.89, 5.41] |
| Subtotal (95% CI) | 147       |    |       | 150       |    |       | 100.0% | 1.99 [-0.43, 4.42] |

Heterogeneity: Chi^2 = 0.07, df = 1 (P = 0.79); P = 0%
Test for overall effect: Z = 1.61 (P = 0.11)

Total (95% CI)

Heterogeneity: Chi^2 = 0.07, df = 1 (P = 0.79); P = 0%
Test for overall effect: Z = 1.61 (P = 0.11)
Test for subarous differences: Not applicable
Comparison of the Boston Carpal Tunnel Questionnaire Symptom Severity Scale (BCTQ-S) score between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.
Comparison of the Boston Carpal Tunnel Questionnaire Functional Status Scale (BCTQ-F) score between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.

Forest plots showing the standardized mean difference for the Semmes-Weinstein (SW) monofilament test between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.
Figure 8

Forest plots showing the standardized mean difference for the Two-point Discrimination (2PD) test between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.

Figure 9

Comparison of overall satisfaction ratings after CTS release between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.
Comparison of key pinch strength between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.

Comparison of the time to return to work (RTW) between patients who underwent ECTR and those who underwent OCTR. OCTR, open carpal tunnel release; ECTR, endoscopic carpal tunnel release; SD, standard deviation; IV, inverse-variance; CI, confidence interval; df, degrees of freedom.
Comparison of all complications between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; M-H, Mantel-Haenszel; CI, confidence interval; df, degrees of freedom.

### Table

| Study or Subgroup       | ECTR Events | ECTR Total | OCTR Events | OCTR Total | Weight | M-H, Fixed, 95% CI | Odds Ratio |
|-------------------------|-------------|------------|-------------|------------|--------|-------------------|------------|
| Agee 1992               | 4           | 82         | 4           | 65         | 10.8%  | 0.78 [0.19, 3.25]  |
| Aslani 2012             | 1           | 32         | 0           | 64         | 0.9%   | 6.14 [0.24, 155.11]|
| Atroshi 2006            | 2           | 63         | 1           | 65         | 2.4%   | 2.16 [0.19, 23.74] |
| Brown 1993              | 4           | 84         | 0           | 85         | 1.2%   | 9.56 [0.51, 180.37]|
| Dumontier 1995          | 2           | 56         | 2           | 40         | 5.6%   | 0.70 [0.09, 5.22]  |
| Ejiri 2012              | 4           | 50         | 0           | 50         | 1.1%   | 9.77 [0.51, 186.52]|
| Erdmann 1994            | 2           | 53         | 7           | 52         | 17.0%  | 0.25 [0.05, 1.28]  |
| Ferdinand 2002          | 1           | 25         | 3           | 25         | 7.2%   | 0.31 [0.03, 3.16]  |
| Gümüştaş 2015           | 1           | 21         | 4           | 20         | 9.7%   | 0.20 [0.02, 1.97]  |
| Jacobsen 1996           | 3           | 18         | 1           | 16         | 2.0%   | 3.46 [0.32, 37.47] |
| Kang 2013               | 0           | 52         | 0           | 52         | Not estimable |
| Larsen 2013             | 6           | 30         | 13          | 60         | 17.3%  | 0.90 [0.31, 2.68]  |
| Macdermid 2003          | 0           | 91         | 0           | 32         | Not estimable |
| Mackenzie 2000          | 0           | 22         | 0           | 14         | Not estimable |
| Martínez-Catasús 2019   | 2           | 35         | 1           | 52         | 1.9%   | 3.09 [0.27, 35.46] |
| Michelotti 2014         | 0           | 30         | 0           | 30         | Not estimable |
| Oh 2017                 | 0           | 35         | 0           | 32         | Not estimable |
| Rab 2006                | 0           | 10         | 0           | 10         | Not estimable |
| Saw 2003                | 3           | 74         | 3           | 76         | 7.1%   | 1.03 [0.20, 5.27]  |
| Sennwald 1995           | 1           | 25         | 2           | 22         | 5.1%   | 0.42 [0.04, 4.94]  |
| Tian 2007               | 3           | 32         | 0           | 34         | 1.1%   | 8.19 [0.41, 165.03]|
| Trumble 2002            | 0           | 97         | 3           | 95         | 8.8%   | 0.14 [0.01, 2.66]  |
| Wong 2003               | 0           | 30         | 0           | 30         | Not estimable |
| Zhang 2016              | 0           | 69         | 0           | 138        | Not estimable |
| Zhao 2004               | 2           | 26         | 0           | 21         | 1.2%   | 4.39 [0.20, 96.54] |
| Total (95% CI)          | 1140        | 1180       | 100.0%      | 100.0%     | 1.06   [0.69, 1.64]|

| Total events            | 41          | 44         |             |           |        |

Heterogeneity: Chi² = 19.00, df = 16 (P = 0.27); I² = 16%

Test for overall effect: Z = 0.28 (P = 0.78)
Comparison of transient nerve injury between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; M-H, Mantel-Haenszel; CI, confidence interval; df, degrees of freedom.
Comparison of permanent nerve injury between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; M-H, Mantel-Haenszel; CI, confidence interval; df, degrees of freedom.
Comparison of scar-related complications between patients who underwent ECTR and those who underwent OCTR. ECTR, endoscopic carpal tunnel release; OCTR, open carpal tunnel release; M-H, Mantel-Haenszel; CI, confidence interval; df, degrees of freedom.
A funnel plot shows the relative symmetry in relation to the pooled estimate from the meta-analysis, indicating no overt publication bias. SE, standard error; OR, odds ratio.

Supplementary Files
This is a list of supplementary files associated with this preprint. Click to download.
PRISMA2009checklist.pdf