



Derkash's Classification and Vas Visual Analog Scale to Access the Long-Term Outcome of Neurothoracic Outlet Syndrome: A Meta-Analysis and Systematic Review

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Specialty section:

This article was submitted to
Headache and Neurogenic Pain,
a section of the journal
Frontiers in Neurology

Received: 18 March 2022

Accepted: 20 May 2022

Published: 06 July 2022

Citation:

Lingyun W, Ke S, Jinmin Z, Yu Q and
Jun Q (2022) Derkash's Classification
and Vas Visual Analog Scale to
Access the Long-Term Outcome of
Neurothoracic Outlet Syndrome: A
Meta-Analysis and Systematic Review.
Front. Neurol. 13:899120.
doi: 10.3389/fneur.2022.899120

Objective: Many publications report outcomes of surgical treatment for neurothoracic outlet syndrome (NTOS); however, high-quality meta-analyses regarding objective evaluation system accessing the long-term outcome of NTOS are lacking. This meta-analysis summarizes and compares the outcomes of Derkash's classification and vas visual analog scale of the supraclavicular neuroplastic of brachial plexus (SNBP) and trams auxiliary first rib resection (TFRR).

Methods: The Cochrane Library, PubMed, EMBASE, Allied and Complementary Medicine (AMED) were searched for papers published between January 1980 and February 2021, using the keywords "thoracic outlet syndrome," "treatment, surgical." Articles were eligible for inclusion if the following criteria were met studies describing outcomes of surgery for NTOS, published in English, human studies, and available in full text. The exclusion criteria were case reports ($n < 10$), reviews, abstracts, and studies lacking a control group or without evaluation for two types of surgery.

Results: We included 10 studies with 1,255 cases, out of which 622 were in the SNBP group; and 633 were in the TFRR group. After surgery (≥ 12 months), Derkash's classification was improved in 425 cases with SNBP and 364 cases with TFRR. OR = 1.34 (95% CI: 0.94, 1.92), $P = 0.03$; vas visual analog scale was improved in 282 cases in the SNBP group and 214 cases in the TFRR group. OR = 1.08 (95% CI: 0.63, 1.85), $P = 0.78$.

Conclusion: This meta-analysis shows that both SNBP and TFRR are effective for NTOS, but that SNBP is better than TFRR in improving Derkash's classification in the long term. Although patients treated with SNBP are more satisfactory, there is no significant difference in vas visual analog scale from TFRR.

Systematic Review Registration: https://www.crd.york.ac.uk/PROSPERO/display_record.asp?ID=CRD42021254203, PROSPERO CRD42021254203.

Keywords: neurologic thoracic outlet syndrome, vas visual analog scale, long-term outcome, meta-analysis, surgical approach (supraclavicular or trams auxiliary)

INTRODUCTION

Neurological thoracic outlet syndrome (NTOS) accounts for ~80%–97% of patients with thoracic outlet syndrome (TOS) (1, 2). Treating priorities with rehabilitation training and analgesia can significantly improve outcomes for most patients (3–5). Surgical treatment may be considered for patients whose conservative treatment was not effective or worsened for more than 6 months (6). Most clinicians believe that surgery has a positive effect on the rapid recovery of the upper limb, pain relief, and low recurrence (7). The surgical treatments mainly include the supraclavicular neuroplastic of brachial plexus (SNBP) and trams auxiliary first rib resection (TFRR). Yin et al. (8) conducted a meta-analysis on the outcome of SNBP and TFRR through 32 studies. The conclusion showed that the SNBP has a high probability of success rate >80% and the TFRR has a high probabilities of success rate >70% but only low probabilities of success rate >80%. However, this review was complicated by lacking quantifiable measures of symptoms and disabilities using validated patient-reported instruments. Previous studies have neglected the access criterion about long-term outcomes. Thus, these results are inadequate for evaluation.

We conducted a meta-analysis with the primary objective of comparing Derkash's classification to access the long-term outcome of NTOS after surgery. Our secondary objective was to compare the vas visual analog scale after surgery and to analyze the reasons for the difference between the two.

MATERIAL AND METHODS

Strategy

This meta-analysis was based on Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines (9). This study has been registered with PROSPERO (https://www.crd.york.ac.uk/PROSPERO/display_record.asp?ID=CRD42021254203).

Literature Search

Two reviewers (W.L.Y and Q.Y) independently conducted a literature search and cross-check. Discussion took place with the third investigator (S.K and Z.J.M) when they had different opinions. We searched the Cochrane Library, PubMed, EMBASE, Allied and Complementary Medicine (AMED) through the network. To avoid missing valid documents, we further searched conference reports and unpublished documents. The keywords: (thoracic outlet syndrome) and (treatment, surgical) were searched from January 1980 to February 2021. Free text words were also used instead of MeSH terms to avoid missing recent publications that were not yet given MeSH headings.

Study Selection

The review identified 3,225 articles. After removal of duplicates and ineligible articles, 647 remained (**Figure 1**). Two authors (W.L.Y and Q.Y) screened the titles and abstracts of the identified studies for relevance. Full texts were obtained of the remaining relevant studies, and two authors (W.L.Y and S.K) read the full-text papers for 175 studies, and a third reviewer (Z.J.M) resolved any disagreements with a final selection. Ten articles that met the inclusion criteria for qualitative analysis were retained.

Data Extraction

Two independent reviewers (W.L.Y and S.K) conducted data extraction. Data on the publication time, male to female incidence rate, follow-up time, data source (country, ethnicity), research institution, surgical approach, research type, and outcomes extracted from the eligible studies. The authors of the original articles were contacted if any discrepancies in data were present. If the opinions of two reviewers were inconsistent, the senior researcher (Z.J.M) was consulted.

Validity Assessment

Two independent reviewers (W.L.Y and S.K) assessed the methodological quality of the articles using the Newcastle-Ottawa Scale (NOS). This scale determines the quality of non-randomized studies based on three categories (selection, exposure and comparability). It allocates zero or 1 point to each numbered item in each category with the exception of comparability where up to 2 points can be assigned. Each study can be allocated a maximum of 9 points, and a score of fewer than 7 points may signify a high risk of bias (10). The NOS was selected because it is validated and adaptable to our meta-analysis.

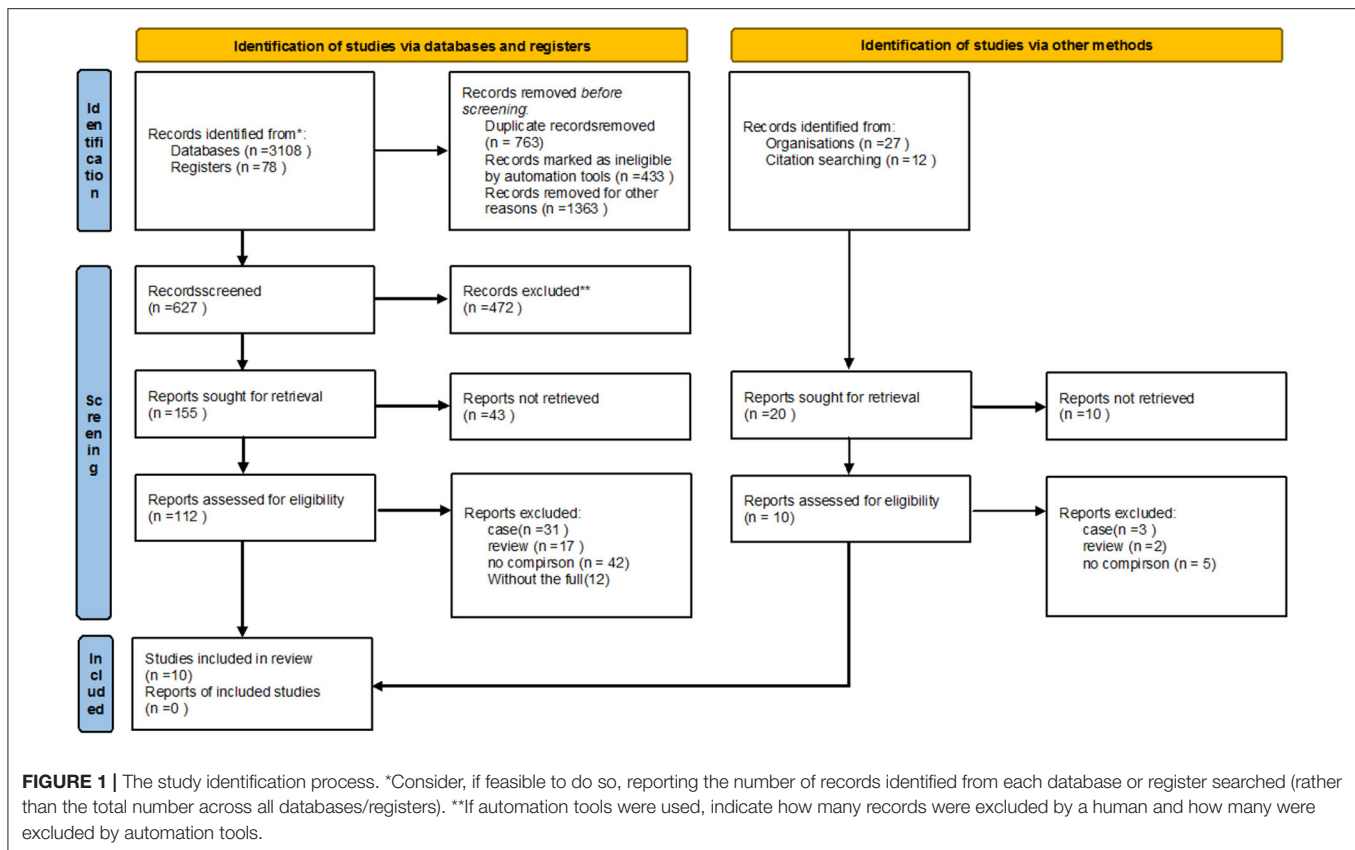
Inclusion and Exclusion Criteria

Types of Studies

The criteria for eligible articles were studies describing outcomes of surgery for NTOS, published in English, human studies, with the full text available. The exclusion criteria were: (1) a review or meta-analysis, (2) consensus, treatment guidelines, (3) case-series and case reports ($N < 10$), (4) studies of endoscopic-assisted or robotic endoscopic-assisted transaxillary first rib resection (TFRR), (5) lacked a control group or were without postoperative follow-up evaluation.

Types of Participants

We included all studies with patients receiving operative interventions for TOS of any type. However, we excluded papers describing patients with compression due to malignancies in athletes and TOS in children (age <12 years). All participants had to be older than 12 years with poor results from conservative treatment or worsening symptoms more than 6 months before



surgery. There was no restriction for sex, socioeconomic status, method of diagnosis, or duration of symptoms, but the follow-up period was set at more than 12 months. The lace was classified according to the definition given by the Oxford Center for Evidence-Based (11). Help from medical professionals who were fluent in both English and the language of the desired publication in times when the authors lacked fluency in the latter. Any inconsistencies within the included works were resolved by contacting the authors of the original studies. Whenever the information could not be obtained, all the reviewers joined the assessment until a consensus was reached.

Outcome Measures

The primary outcome measures were results of surgical treatment that were reported using validated questionnaires and outcomes according to Derkash's classification or similar. It is divided into excellent result (E): no pain, easy return to preoperative professional and leisure daily activities; good result; (G): intermittent pain well tolerated, possible return to preoperative professional and leisure daily activities; fair result (F) intermittent or permanent pain with bad tolerance, difficult return to preoperative professional and daily activities; poor result (P): symptoms not improved or aggravated. We extracted excellent and good as improved function, fair and poor as no improvement.

The secondary outcome was relieved in pain at least 12 months after the intervention preferably measured as change on a validated visual analog scale (VAS) or similar.

Definition

Long term outcome: the outcome about evaluation for NTOS after surgery (> 12 months).

Short term outcome: the outcome about evaluation for NTOS after surgery (1 week to 3 months).

Derkash's classification: it is used to assess perceived disability in patients with arm, shoulder, and hand problems. These assessment schemes can well reflect the function of the upper limb and categorize primary outcomes as "excellent," "good," "fair," and "poor".

Statistical Analysis

Statistical analysis was conducted by two reviewers (W.L.Y and S.K). We used Revman5.3 by Cochrane. A random-effects model was used for pain relieve with validated visual analog scale (VAS) and fix-effects model was used for improvement of upper limb with Derkash's classification. While probing for heterogeneity between the included studies, the Chi² test and the Higgins I² statistic were used. Cochran's Q P-value of <0.10 for the Chi² test was chosen to show significant heterogeneity between the studies. Higgins I² values between 0 and 40% were considered "might not be important," 30%–60% as "might show moderate

TABLE 1 | The characteristics of included studies.

Study	Country	Type of study	Number of patients	Average-age (year)	Outcome	Follow-up (month)
Aboul Hosn (14)	USA	Retros-pective	82	32.2	Upper limb function (Derkash's classification) and Complication: 1. In the SNBP group, 60 out of 63 had improved upper limb function. Twenty-one out of 32 in TFRR improved upper limb function. 2. There were 30 cases of complications after SNBP, 25 cases of pneumothorax, 3 cases of hemothorax, and 2 cases of vascular injury. Fifteen cases of postoperative complications of TFRR, 11 cases of pneumothorax, 2 cases of hemothorax, 2 cases of hemopneumothorax, 1 case of vascular injury	17.9
Altobelli (15)	USA	Retros-pective	334	40	Vas score (visual analog scale) and upper limb function (Quick-DASH score): 1. The satisfaction rate of TFRR dropped to 45% (254 cases) at 24 months and to 38% (118 cases) after 36 months. 2. The satisfaction rate of SNBP was 80% (80 cases) at 12 months, 69% at 24 months, and 59% (45 cases) at 36 months. The difference were statistically significant ($P < 0.05$). The recurrence rate is higher (54%) in TFRR	25
Balci (16)	Turkey	Retros-pective	47	37.9	The changes of ulnar nerve conduction velocities (UNCV); complications; mortality; upper limb function (Derkash's classification): The results showed that there was no significant difference between the TFRR and SNBP	55
Bhattacharya (17)	UK	Retros-pective	70	37	Pain relieve (visual analog scale) and upper limb function (Derkash's classification): The results showed that postoperative pain was relieved and upper limb function was improved in both TFRR and SNBP	43
Cikrit (18)	USA	Retros-pective	37	37.5	Blood loss; complications; upper limb function (Cervical Brachial Score Questionnaire): The results showed that SNBP had less blood loss (61 vs. 218 cc), fewer complications (1 vs. 21), and higher improvement in upper limb function (100 vs. 83%) than TFRR	36
Degeorges (19)	France	Retros-pective	176	35.7	Complications and improvement of upper limb function (Derkash's classification): The final follow-up was 69 cases in the SNBP group and 107 TFRR cases. The results showed that upper limb function improved in 43 cases in SNBP and 49 cases in TFRR	90
Nasim (20)	USA	Retros-pective	34	37	The upper limb function (Derkash's classification): Among the follow-up cases, 24 in SNBP and 13 in TFRR. The results showed that 20 cases in the SNBP improved, seven cases in the TFRR improved	> 12
Parry (21)	England	Retros-pective	26	36.2	Upper limb function (Quick-DASH score): Among the cases, 13 in SNBP and 12 in TFRR. The results showed that upper limb function improved in 11 cases in SNBP and 8 cases in TFRR	>12
Sanders (22)	USA	Retros-pective	491	34	Pain relieve (visual analog scale) and upper limb function (Quick-DASH): Among the follow-up cases, there were 279 cases in the SNBP and 111 cases in the TFRR. After SNBP, 173 cases of pain were relieved and upper limb function were improved; 66 cases of TFRR achieved the same results, and there was no statistical difference in pain relief and improvement of upper limb function between the two	24
Sheth (13)	USA	Randomized -study	47	37	Vas score (visual analog scale): There was a statistically significant difference between TFRR and SNBP ($P = 0.03$)	47

heterogeneity," 50%–90% as "may show large heterogeneity," and 75%–100% as "may represent notable heterogeneity" (12).

Publication Bias

We checked for publication bias by building a funnel plot to visually check for asymmetry.

RESULTS

Study Identification

The study identification terminated the plan, and 3,225 records were found. According to the exclusion and inclusion criteria ($n = 763$), articles ($n = 433$), review articles ($n = 472$) and articles

published without a control group ($n = 1,363$) were excluded. A total of 10 studies were included in this meta-analysis.

Characteristics of Included Studies

The characteristics of the included studies were presented in **Table 1**. One was a randomized controlled trial (13), nine were prospective controlled observational studies (14–22). To evaluate the long-term outcome of patients with NTOS, other types of TOS were excluded. Finally, 1,255 cases were included, including 622 in the SNBP group and 633 in the TFRR group. Four studies (13, 15, 17, 22) described pain relief with improvement of visual analog scale. Finally, 843 cases were included, including 440 in the SNBP group and 403 in the TFRR group. There was no difference in age, sex ratio, or the number of people in the two groups ($P > 0.05$).

Altobelli et al. (15) conducted a retrospective analysis of the postoperative effects of 254 CASES of NTOS. Vas visual analog Scale and Derkash's Classification were used to compare the improvement of upper limb function and pain relief after surgery, and the results showed that the satisfaction rate of TFRR was 53% at 12 months after surgery and decreased to 45% at 24 months and 38% at 36 months. The satisfaction rate was 80% at 12 months, 69% at 24 months, and 59% at 36 months after SNBP surgery. There were significant differences in upper limb function improvement and pain relief between the two methods ($P < 0.05$). In addition, the TFRR group had a higher postoperative recurrence rate (54%), requiring SNBP treatment again.

Aboul Hosn et al. (14) conducted a retrospective study on the postoperative follow-up results of 95 CASES of NTOS, including 63 cases of SNBP and 32 cases of TFRR. The follow-up time was over 24 months. Derkash's classification was used to evaluate the postoperative upper limb function, and the incidence of complications was compared between the two. The results showed that there were 30 complications after SNBP, including pneumothorax in 25 cases, hemothorax in three cases and vascular injury in two cases. Postoperative complications of TFRR included 15 cases, 11 cases pneumothorax, two cases hemothorax, two cases hemopneumothorax, and one case vascular injury. Postoperative upper limb function improved in 60 cases with SNBP and 21 cases with TFRR.

Bhattacharya et al. (17) retrospectively observed 60 patients after NTOS using the Derkash's Classification scale questionnaire, with a follow-up rate of 90% and a mean follow-up period of 43 months. The results showed that postoperative pain was relieved in TFRR and SNBP groups, and upper limb function was improved to some extent. There was no difference in long-term results between the two methods, and postoperative complications were not clearly described.

Sanders et al. (22) reviewed 491 NTOS patients with follow-up >24 months for postoperative evaluation. There were 279 cases in the SNBP group and 111 cases in the TFRR group. The upper limb pain was relieved and the function was improved in 173 cases after SNBP. The same results were obtained in 66 cases of TFRR, and there was no statistical difference in pain relief and upper limb function improvement between the two methods.

Sheth et al. (13) was the only RCT article in which the included patients were divided into two groups according to a completely

TABLE 2 | The Newcastle-Ottawa Scale score for studies.

Qua Quality assessment of Newcastle-Ottawa Scale score by Assessment of the Studies Using Newcastle-Ottawa Scale for Case-Control Studies

Study number	First author	Case definition adequate	Representativeness of cases	Selection of controls	Definition of controls	Comparability	Ascertainment of exposure	Same ascertainment method	Nonresponse rate	Total
1	Altobelli (15)	*	*	*	*	*	*	*	*	8
2	Aboul Hosn (14)	*	*	*	*	*	*	*	*	7
3	Balci (16)	*	*	*	*	**	*	*	*	9
4	Bhattacharya (17)	*	*	*	*	*	*	*	*	6
5	Clkrit (18)	*	*	*	*	**	*	*	*	9
6	Degeorges (19)	*	*	*	*	*	*	*	*	8
7	Nasim (20)	*	*	*	*	*	*	*	*	8
8	Parry (21)	*	*	*	*	*	*	*	*	7
9	Sanders (22)	*	*	*	*	*	*	*	*	8
10	Sheth (13)	*	*	*	*	**	*	*	*	9

A maximum of one "star" for each item within the "Selection" and "Exposure/Outcome" categories; maximum of two "stars" for "Comparability."

randomized control, with 24 patients in the TFRR group and 25 patients in the SNBP group. The mean follow-up time was 37 months, and the postoperative efficacy was evaluated by VAS. Results Preoperative VAS scores were 77 ± 3 in TFRR group and 82 ± 3 in SNBP group, with no difference between the two groups ($P = 0.28$). Postoperative VAS scores of TFRR group (39 ± 7) and SNBP group (61 ± 7) were significantly different ($P = 0.03$).

Test of Heterogeneity

We performed statistical testing for heterogeneity to decide if the long-term outcomes of surgical treatment were the same in the included studies. The Cochran Q result showed that there was mild heterogeneity between studies ($\chi^2 = 11.90, P = 0.22$). In addition, I^2 revealed that 24% of the variation across the studies was because of heterogeneity rather than sampling error and chance.

Quality Assessment

The mean Newcastle-Ottawa Scale score for 10 retrospective studies was 8.0, with higher scores showing better quality (Table 2).

Improvement of Derkash’s Classification

Ten studies ($n = 1,255$ cases) were included, 622 in the SNBP group and 633 in the TFRR group. According to Derkash’s classification, improvements in the upper limb were achieved in 425 cases with SNBP and 364 cases with TFRR. The excellent and good rate of SNBP was 1.34 times that of TFRR. $OR = 1.34$ (95% CI: 0.94, 1.92), $P = 0.03$ (Figure 2).

Improvement of Vas Visual Analog Scale (VAS Score)

Based on the random-effects model for four studies with the changes of VAS. We note that 282 patients in the SNBP group had significant relief of pain and that there was no need to continue taking analgesics or reduce the dose for these patients; 214 patients in the TFRR group gained similar results. There was no difference between the two. $OR = 1.08$ (95% CI: 0.63, 1.85), $P = 0.78$ (Figure 3).

Publication Bias

We set up a funnel plot to assess publication bias. In the non-publication bias, we would expect the studies to be symmetrical about the combined effect size. The remarked asymmetry in our funnel plot suggested the likelihood of some publication bias (Figure 4). This was mostly true when the analysis included clinical trials to find bias in publishing null results. However, regarding our research, this meant that including better-powered studies might have led to low-bias meta-analysis. The publication bias section and the idea of having a funnel plot was a technique to highlight any statistically probable error in the review, aiming to raise the truth of the conclusion drawn.

DISCUSSION

Importance of Long-Term Outcome Assessment

Neurothoracic outlet syndrome has been perplexing clinicians for effective treatment because of the lack of Consistent and objective examinations (23). For patients who have been treated

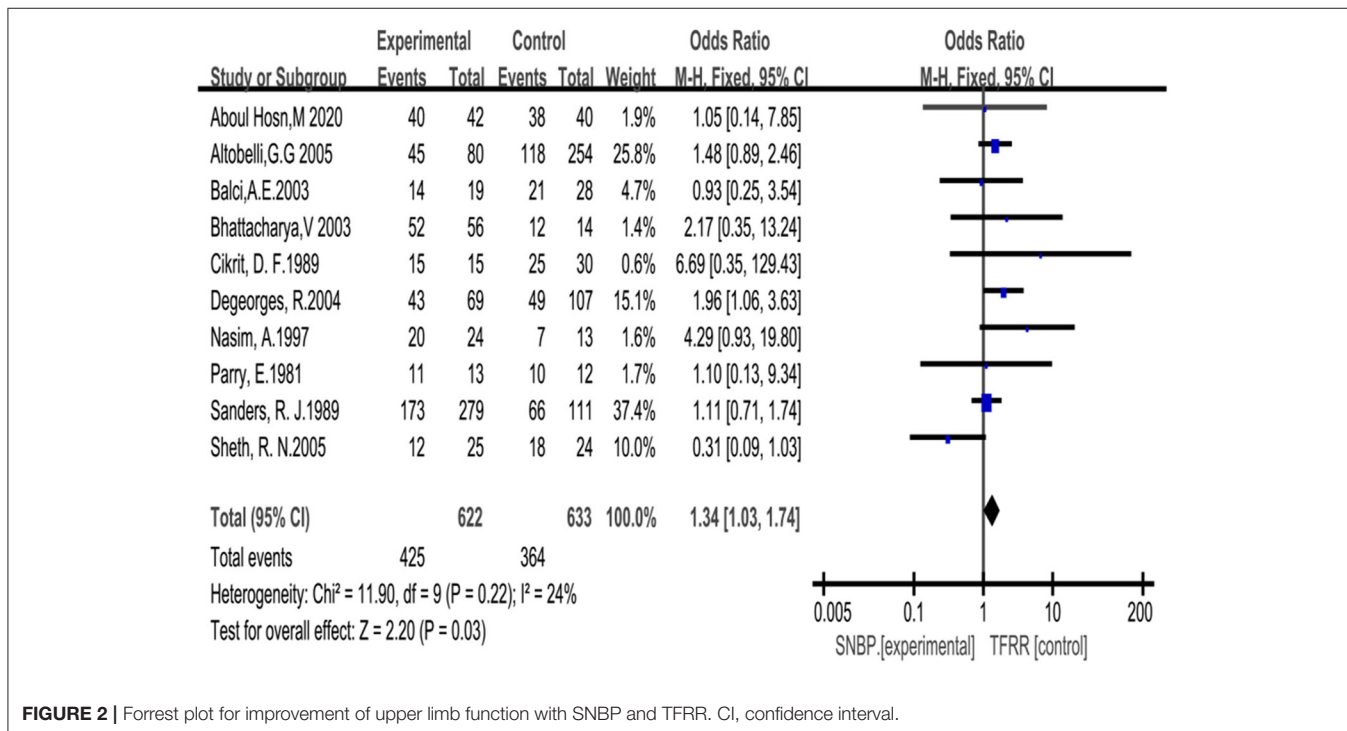


FIGURE 2 | Forrest plot for improvement of upper limb function with SNBP and TFRR. CI, confidence interval.

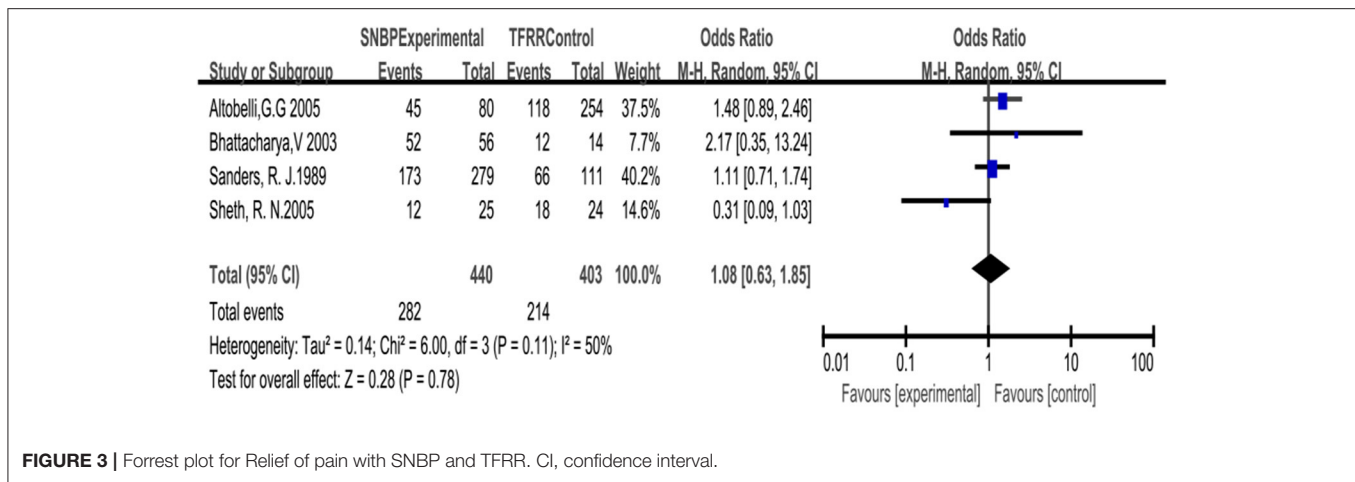


FIGURE 3 | Forrest plot for Relief of pain with SNBP and TFRR. CI, confidence interval.

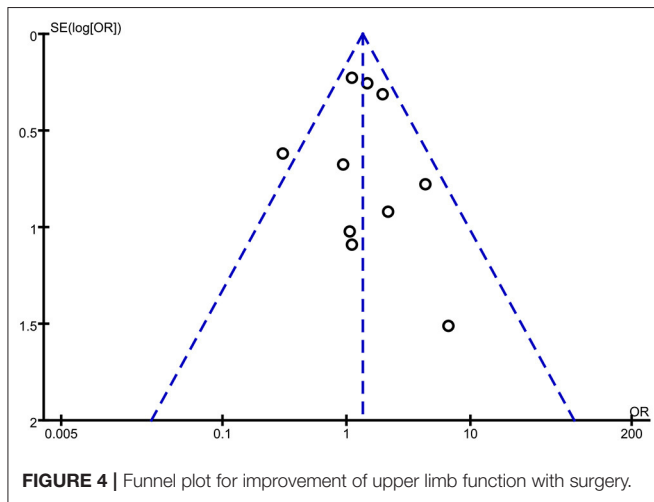


FIGURE 4 | Funnel plot for improvement of upper limb function with surgery.

conservatively but ineffectually for more than 6 months, most clinicians prefer surgical treatment. Certainly, the long-term outcome of surgery directly affects the evaluation for treatment, as it is more instructive and convincing than the short-term outcome. Although many patients have good short-term results in their upper limb after surgery, over time, the results will gradually worsen or recur. Yoshizumi et al. (24) believed that long-term evaluation after treatment can be more stable and objective. However, choosing the better surgery has always been controversial (16, 25). This study aimed to systematically analyze the studies and provide the most comprehensive data thus far on the long-term results of surgery to determine the difference.

Improvement of Derkash's Classification

Derkash's classification is a region-specific tool, as it assesses perceived disability in patients with arm, shoulder, and hand problems. These assessment schemes can well reflect the function of the upper limb and are used to categorize primary outcomes as "excellent" if there was no pain and a return to preoperative status, "good" if intermittent pain was well tolerated and there

was the possibility of a return to the preoperative status, "fair" if intermittent or permanent pain was not well tolerated and a return to the preoperative status was difficult, or "poor" if symptoms had not improved or were aggravated (26). Peek et al. (27) confirmed that Derkash's classification was an effective tool to evaluate the improvement of upper limb function because of its wide use in practice and the availability of the data. Ruopsa et al. (28) conducted a survey lasting for 12 years that included 94 NTOS patients evaluated through Derkash's classification, The results showed that most patients had significant improvement by SNBP. He believed that resection of the first rib was unnecessary. Johansen et al. (29) confirmed this through more cases (504 cases). We included 1,255 cases in the study. The excellent and good rate of SNBP was 1.34 times that of TFRR. OR = 1.34 (95% CI: 0.94, 1.92), $P = 0.03$.

Improvement of Vas Visual Analog Scale (VAS Score)

The VAS score is widely used in pain assessment and is accurate, simple, and highly sensitive (30). Sheth et al. (13) divided patients into two groups according to a randomized control. The results showed that postoperative VAS scores in the TFRR (39 ± 7) and SNBP (61 ± 7) groups were significantly different ($P = 0.03$). However, this study had selection bias and few cases (45 cases), so it could not be concluded that TFRR was better than SNBP in relief pain. We included 843 cases in the study, the cases were enormous, there was no significant difference between the two.

The Difference Between TFRR and SNBP

In the past, most doctors believed the occurrence of NTOS was related to compression of the brachial plexus by the first rib (31). Therefore many surgical approaches for removing the first rib, especially complete or partial removal of the first rib became popular (32, 33). TFRR achieved good results (48%), but the recurrence was still high (60%–70%) and often caused damage to the vessel and brachial plexus (34). Some studies showed that the NTOS was more likely to be related to the entrapment of the scalene muscles and bands on the brachial plexus. SNBP exposed the brachial plexus more fully, removed the scalene muscles

and abnormal bands more thoroughly, and the effect was more precise than TFRR (35). However, scholars who agreed that the occurrence of NTOS stemmed from the first rib believed that TFRR could be more suited for skeletal compression (36).

Knowledge of the anatomy of the brachial plexus and thoracic outlet revealed that hypertrophy of the scalene muscle, fibrosis, and band compression were essential causes of neurovascular compression (37). Therefore, it was difficult to release the compression where the C5-T1 rami roots passed through the intervertebral foramen through TFRR, which was the main reason that the compression couldn't be relieved. SNBP can more fully disclose the C5-T1 roots by removing part of the scalene muscles, especially the portion where the brachial plexus penetrates the intervertebral foramen. The intraoperative release was more complete, and there was no need to remove the first rib, which avoided neurovascular damage and safer.

Our study does have several limitations. First, most studies were retrospective, only 1 randomized controlled trial was identified. The included studies are of moderate to poor methodological quality, as assessed using the NOS score; therefore, the overall quality of the available evidence is low, and there is a high risk of bias. Our analysis relied on these reported data, but we cannot improve the quality of the data. Second, the lack of unified evaluation criteria for NTOS creates a risk of bias. There are many evaluation criteria for improvement in upper limb, such as Derkash's Classification, Quick Shoulder and Hand questionnaire (Quick-dash Score) and Cervical Brachial Score Questionnaire (CBSQ). Derkash's classification was selected as the primary outcome measure in our analysis, other evaluation systems were excluded. Therefore, perhaps some interesting studies might have been ignored in this review. However, we listed all excluded studies with their characteristics and reason for exclusion. Third, this meta-analysis was limited

because of the lack of objective evaluation criteria among the included studies. Although we conducted a comparison with ulnar nerve conduction velocities (UNCV), data on this topic are rare.

Finally, our results focus on patient reports of improvement in upper limb. Although VAS Score was used to analyze pain relief, only a few studies had objective data. Comparison using more objective evaluation criteria such as UNCV changes and VAS Score will make outcome more convincing. In addition, retrospective studies lack randomized controls, which has a high risk of bias. Multicenter, randomized controlled studies are needed in the future.

In conclusion, this meta-analysis shows that both SNBP and TFRR are effective for NTOS, but that SNBP is better than TFRR in improving Derkash's classification in the long term. Although patients treated with SNBP were more satisfactory, there was no significant difference in vas visual analog scale from TFRR.

AUTHOR CONTRIBUTIONS

WL is responsible for literature collection, data analysis, and writing articles. ZJ and WL was in charge of experimental design. ZJ and SK was responsible for evaluation of literature quality. SK was responsible for analyzing the data. QY and QJ are responsible for literature search and data collection. All authors contributed to the article and approved the submitted version.

FUNDING

This work was funded by Science and Technology Research Program (1598011-5) and Science and Technology Department of Guangxi Zhuang Autonomous Region.

REFERENCES

- Masocatto NO, Da-Matta T, Prozzo TG, Couto WJ, Porfirio G. Thoracic outlet syndrome: a narrative review. *Rev Col Bras Cir.* (2019) 46:e20192243. doi: 10.1590/0100-6991e-20192243
- Kaplan J, Kanwal A. *Thoracic Outlet Syndrome*, in *StatPearls*. StatPearls Publishing Copyright © 2021. Treasure Island, FL: StatPearls Publishing LLC (2021).
- Ferrante MA, Ferrante ND. The thoracic outlet syndromes: part 2. The arterial, venous, neurovascular, and disputed thoracic outlet syndromes. *Muscle Nerve.* (2017) 56:663–73. doi: 10.1002/mus.25535
- Pesser N, Goeteyn J, van der Sanden L, Houterman S, van Alfen N, van Sambeek MRHM. Feasibility and outcomes of a multidisciplinary care pathway for neurogenic thoracic outlet syndrome: a prospective observational cohort study. *Eur J Vasc Endovasc Surg.* (2021) 61:1017–24. doi: 10.1016/j.ejvs.2021.02.048
- Goeteyn J, van den Borek R, Bouwman A, Pesser N, van Nuenen B, van Sambeek M. Interfascial plane blocks reduce postoperative pain and morphine consumption in thoracic outlet decompression. *Ann Vasc Surg.* (2020) 66:301–8. doi: 10.1016/j.avsg.2019.12.005
- Boglione M, Ortiz R, Teplisky D, Giuseppucci C, Korman L, Reusmann A, Barrenechea M. Surgical treatment of thoracic outlet syndrome in peditriacs. *J Pediatr Surg.* (2021) 4:581–9. doi: 10.1016/j.jpedsurg.2021.08.017
- Franklin GM. Work-related neurogenic thoracic outlet syndrome: diagnosis and treatment. *Phys Med Rehabil Clin N Am.* (2015) 26:551–61. doi: 10.1016/j.pmr.2015.04.004
- Yin ZG, Gong KT, Zhang JB. Outcomes of surgical management of neurogenic thoracic outlet syndrome: a systematic review and bayesian perspective. *J Hand Surg Am.* (2019) 44:416.e1–17. doi: 10.1016/j.jhsa.2018.06.120
- Liberati A, Altman DG, Tetzlaff J, Mulrow C, Gotzsche PC, Ioannidis JPA. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. *PLoS Med.* (2009) 6:e1000100. doi: 10.1371/journal.pmed.1000100
- Stang A. Critical evaluation of the Newcastle-Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol.* (2010) 25:603–5. doi: 10.1007/s10654-010-9491-z
- Atkins D, Eccles M, Flottorp S, Guyatt GH, Henry D, Hill S. Systems for grading the quality of evidence and the strength of recommendations I: critical appraisal of existing approaches The GRADE Working Group. *BMC Health Serv Res.* (2004) 4:38. doi: 10.1186/1472-6963-4-38
- Henry BM, Tomaszewski KA, Walocha JA. Methods of evidence-based anatomy: a guide to conducting systematic reviews and meta-analysis of anatomical studies. *Ann Anat.* (2016) 205:16–21. doi: 10.1016/j.aanat.2015.12.002
- Sheth RN, Campbell JN. Surgical treatment of thoracic outlet syndrome: a randomized trial comparing two operations. *J Neurosurg Spine.* (2005) 3:355–63. doi: 10.3171/spi.2005.3.5.0355

14. Aboul Hosn M, Goffredo P, Man J, Nicholson R, Kresowik T, Sharafuddin M. Supraclavicular versus transaxillary first rib resection for thoracic outlet syndrome. *J Laparoendosc Adv Surg Tech A*. (2020) 30:737–41. doi: 10.1089/lap.2019.0722
15. Altobelli GG, Kudo T, Haas BT, Chandra FA, Moy JL, Ahn SS. Thoracic outlet syndrome: pattern of clinical success after operative decompression. *J Vasc Surg*. (2005) 42:122–8. doi: 10.1016/j.jvs.2005.03.029
16. Balci AE, Balci TA, Cakir Ö, Eren S, Eren MN. Surgical treatment of thoracic outlet syndrome: effect and results of surgery. *Ann Thorac Surg*. (2003) 75:1091–6. discussion 1096. doi: 10.1016/S0003-4975(02)04725-2
17. Bhattacharya V, Hansrani M, Wyatt MG, Lambert D, Jones NAG. Outcome following surgery for thoracic outlet syndrome. *Eur J Vasc Endovasc Surg*. (2003) 26:170–5. doi: 10.1053/ejvs.2002.1891
18. Cikrit DF, Haefner R, Nichols WK, Silver D. Transaxillary or supraclavicular decompression for the thoracic outlet syndrome. A comparison of the risks and benefits. *Am Surg*. (1989) 55:347–52.
19. Degeorges R, Reynaud C, Becquemain JP. Thoracic outlet syndrome surgery: long-term functional results. *Ann Vasc Surg*. (2004) 18:558–65. doi: 10.1007/s10016-004-0078-6
20. Nasim A, Sayers RD, Healey PA, Bell PR, Barrie WW. Surgical decompression of thoracic outlet syndrome; is it a worthwhile procedure? *J R Coll Surg Edinb*. (1997) 42:319–23.
21. Parry E. The thoracic outlet compression syndrome. *Aust N Z J Surg*. (1981) 51:84–91. doi: 10.1111/j.1445-2197.1981.tb05915.x
22. Sanders RJ, Pearce WH. The treatment of thoracic outlet syndrome: a comparison of different operations. *J Vasc Surg*. (1989) 10:626–34. doi: 10.1016/0741-5214(89)90005-0
23. Ortaç EA, Sarpel TC, Benlidayi I. Effects of Kinesio Taping on pain, paresthesia, functional status, and overall health status in patients with symptomatic thoracic outlet syndrome: a single-blind, randomized, placebo-controlled study. *Acta Orthop Traumatol Turc*. (2020) 54:394–401. doi: 10.5152/j.aott.2020.19042
24. Yoshizumi T, Murata H, Kanno H, Shinonaga M. Efficacy of supraclavicular scalenotomy followed by external neurolysis without rib resection for post-traumatic neurogenic thoracic outlet syndrome. *Spine*. (2021) 46:E632–8. doi: 10.1097/BRS.0000000000003859
25. Patetta MJ, Naami E, Sullivan BM, Gonzalez MH. Nerve compression syndromes of the shoulder. *J Hand Surg Am*. (2021) 46:320–6. doi: 10.1016/j.jhsa.2020.09.022
26. Al Rstum Z, Tanaka A, Sandhu HK, Miller CC 3rd, Saqib NU, Besho JM, et al., Differences in quality of life outcomes after paraclavicular decompression for thoracic outlet syndrome. *J Vasc Surg*. (2020) 72:1421–6. doi: 10.1016/j.jvs.2019.12.037
27. Peek J, Vos CG, Ünlü Ç, van de Pavoordt HDWM, van den Akker PJ, de Vries JPM. Outcome of surgical treatment for thoracic outlet syndrome: systematic review and meta-analysis. *Ann Vasc Surg*. (2017) 40:303–26. doi: 10.1016/j.avsg.2016.07.065
28. Ruopsa N, Ristolainen L, Vastamäki M, Vastamäki H. Neurogenic thoracic outlet syndrome with supraclavicular release: Long-term outcome without rib resection. *Diagnostics (Basel)*. (2021) 11:450. doi: 10.3390/diagnostics11030450
29. Johansen K. Rib-sparing scalenectomy for neurogenic thoracic outlet syndrome: Early results. *J Vasc Surg*. (2021) 73:2059–63. doi: 10.1016/j.jvs.2020.12.052
30. Karcioglu O, Topacoglu H, Dikme O, Dikme O. A systematic review of the pain scales in adults: which to use? *Am J Emerg Med*. (2018) 36:707–14. doi: 10.1016/j.ajem.2018.01.008
31. Ferrante MA, Ferrante ND. The thoracic outlet syndromes: part 1. Overview of the thoracic outlet syndromes and review of true neurogenic thoracic outlet syndrome. *Muscle Nerve*. (2017) 55:782–93. doi: 10.1002/mus.25536
32. Mingoli A, Feldhaus RJ, Farino C, Cavallari N, Sapienza P, di Marzo L. Long-term outcome after transaxillary approach for thoracic outlet syndrome. *Surgery*. (1995) 118:840–4. doi: 10.1016/S0039-6060(05)80274-0
33. Kawada S, Suzuki I, Inamura S, Ogawa J, Inoue K, Koide S. First rib resection in the treatment of the thoracic outlet syndrome. *Tokai J Exp Clin Med*. (1981) 6:285–95.
34. Seifert S, Sebesta P, Klenske M, Esche M. [Thoracic outlet syndrome]. *Zentralbl Chir*. (2017) 142:104–12. doi: 10.1055/s-0042-121611
35. Caputo FJ, Wittenberg AM, Vemuri C, Driskill MR, Earley JA, Rastogi R. Supraclavicular decompression for neurogenic thoracic outlet syndrome in adolescent and adult populations. *J Vasc Surg*. (2013) 57:149–57. doi: 10.1016/j.jvs.2012.07.025
36. Kocher GJ, Zehnder A, Lutz JA, Schmidli J, Schmid RA. First rib resection for thoracic outlet syndrome: the robotic approach. *World J Surg*. (2018) 42:3250–5. doi: 10.1007/s00268-018-4636-4
37. Machleder HI, Moll F, Verity MA. The anterior scalene muscle in thoracic outlet compression syndrome. Histochemical and morphometric studies. *Arch Surg*. (1986) 121:1141–4. doi: 10.1001/archsurg.1986.014001004700

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