

Contents lists available at [ScienceDirect](#)

## Journal of Hand Surgery Global Online

journal homepage: [www.JHSGO.org](http://www.JHSGO.org)

## Original Research

## Management of Acute Carpal Tunnel Syndrome: A Systematic Review

Ying C. Ku, BS, <sup>\*</sup>, <sup>†</sup> Megan Gannon, DO, <sup>†</sup> Wei Fang, PhD, <sup>‡</sup> Rebecca C. Norcini, MD, <sup>\*</sup>  
Kerri M. Woodberry, MD, MBA <sup>\*</sup><sup>\*</sup> Department of Plastic and Reconstructive Surgery, West Virginia University School of Medicine, Morgantown, WV<sup>†</sup> Campbell University School of Osteopathic Medicine, Lillington, NC<sup>‡</sup> West Virginia Clinical and Translational Science Institute, West Virginia University Health Sciences Center Erma Byrd Biomedical Research Center, Morgantown, WV

## ARTICLE INFO

## Article history:

Received for publication March 6, 2023

Accepted in revised form June 10, 2023

Available online July 20, 2023

## Key words:

acute carpal tunnel syndrome  
carpal tunnel release  
hand surgery  
nerve compression  
nonoperational  
operational  
splinting**Purpose:** This review aims to compare recovery outcomes of conservative, early operative, and a combination of conservative and operative management for acute carpal tunnel syndrome (ACTS).**Methods:** A literature search of PubMed, Scopus, and CINAHL from 1970 to 2022 was conducted using the keywords carpal tunnel syndrome and acute nerve compression. ACTS was defined as a case within 12 weeks of symptom onset. Primary data extracted included causes (traumatic or atraumatic), symptom duration (<1 day, 1–7 days, or 8–84 days), intervention (surgical, conservative, or conservative then surgical), follow-up duration, and outcome (full recovery or non-full recovery). Logistic regression analyses and  $\chi^2$  tests were performed to investigate associations among these variables.**Results:** A total of 197 patients involving 127 (64.5%) traumatic and 70 (35.3%) atraumatic cases were included. Forty-seven percent of patients were managed conservatively followed by surgery, 30% conservative only, and 23% surgery only. The traumatic group was associated with better recovery than the atraumatic group. Recovery outcomes were not associated with symptom duration or follow-up time. The choice of intervention was not associated with traumatic or atraumatic etiology, nor did it affect recovery outcomes in either group.**Conclusions:** Traumatic ACTS is associated with better recovery outcomes than atraumatic etiologies. Surgical intervention was not found to be associated with better outcomes than conservative management, regardless of the etiologies. Further prospective studies are warranted to compare surgical versus conservative management.**Clinical Relevance:** Currently, there are no guidelines for the best management of ACTS, and it is not known if early or delayed surgical treatment is optimal. This review compiles the current evidence and identifies gaps in the literature, highlighting the need for further investigation to provide the best clinical practice.Copyright © 2023, THE AUTHORS. Published by Elsevier Inc. on behalf of The American Society for Surgery of the Hand.  
This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Carpal tunnel syndrome is a well-known neuropathy resulting from median nerve compression under the transverse carpal ligament, leading to symptoms such as pain, numbness, tingling, and weakness in the median nerve distribution of the hand. Carpal tunnel syndrome can be acute or chronic, with a chronic progressive course that lasts for at least 3 months being the most

common.<sup>1–4</sup> Although most cases can present with mild intermittent symptoms, permanent deficits can develop in severe cases if left untreated.<sup>1,5</sup>

Acute carpal tunnel syndrome (ACTS) is a rare form of the carpal tunnel spectrum, characterized by its rapid progressive course and high potential for permanent changes in the local structures including nerves and muscles.<sup>6</sup> The onset of symptoms for ACTS typically ranges from hours to days following an inciting event, with symptoms similar to the chronic form except for the acute onset and severity tending to be more debilitating. It is most commonly associated with wrist trauma and was reported to be a complication secondary to local hemorrhage and edema in 4.3% of distal radius fractures.<sup>7</sup> Besides traumatically induced injuries,

**Declaration of interests:** No benefits in any form have been received or will be received related directly to this article.

**Corresponding author:** Kerri M. Woodberry, MD, MBA, West Virginia University School of Medicine, 1 Medical Center Drive PO Box 9238, Morgantown, WV 26506.

E-mail address: [kerri.woodberry@hsc.wvu.edu](mailto:kerri.woodberry@hsc.wvu.edu) (K.M. Woodberry).

<https://doi.org/10.1016/j.jhsg.2023.06.012>

2589-5141/Copyright © 2023, THE AUTHORS. Published by Elsevier Inc. on behalf of The American Society for Surgery of the Hand. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

atraumatic causes have also been identified including hemorrhagic disorders, rheumatologic disorders, vascular disorders, infections, and burns.<sup>5,6,8</sup> Regardless of the inciting events, a rapid increase in intracompartmental pressure secondary to a mass effect is thought to be the common pathway leading to ACTS. Once the pressure exceeds a certain threshold, local blood flow is compromised and ischemia with secondary nerve conduction dysfunction and endoneurial edema can develop.<sup>8,9</sup> Several studies have reported the prevalence of carpal tunnel syndrome in the adult population ranging from 1% to 5% depending on diagnostic guidelines with a higher prevalence associated with obesity, older age, and woman.<sup>10,11</sup> However, there is limited representative data on the incidence and prevalence of ACTS. Nevertheless, ACTS is an important disease given its potential to cause permanent damage to the hand.

Although surgical intervention is reported as the preferred treatment in some traumatic ACTS cases,<sup>12</sup> others have reported successful treatment through conservative management.<sup>3,13</sup> There are no established guidelines or algorithms to direct treatment approaches. Whether to first approach ACTS with conservative management remains unclear. Therefore, the objective of this review is to investigate current evidence in the literature regarding management preferences and to compare different treatment approaches and their outcomes.

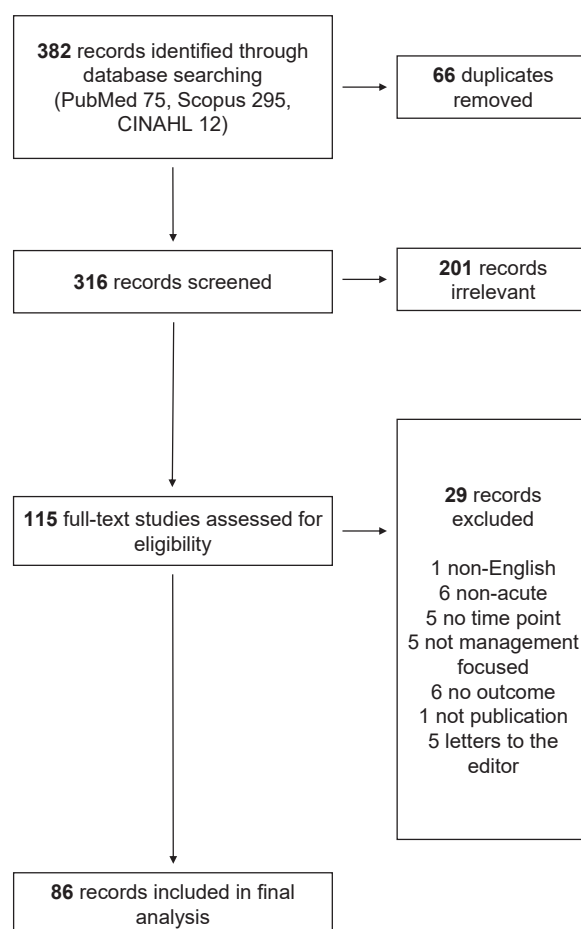
## Methods

### Search strategy and study selection

A comprehensive literature search of PubMed, Scopus, and CINAHL was conducted in April 2022 to identify studies that reported ACTS management in the adult population (18+ years). Literature was limited to those written in English and published from 1970 to 2022. Medical Subject Heading (MeSH) terms and keywords including Carpal Tunnel Syndrome [MeSH], Acute Disease [MeSH], and adults were used to identify literature of interest. Following duplicates removal, all titles and abstracts identified by the search criteria were screened independently by two authors (Y.K. and M.G.) for relevance. Full-text review of the remaining articles for eligibility was conducted by the same two authors independently. Carpal tunnel syndrome cases were determined as acute when patients presented to a health care setting within 12 weeks after the onset of symptoms.<sup>2–4</sup> Articles were excluded if they met the following: were not written in English, were not in an acute setting, did not provide acute timepoint, did not discuss treatment/management, were not in the adult population, or had no outcome assessment following treatment. Articles without a publication and letters to the editor were also excluded. A consensus was reached through discussion between the above two authors in cases of disagreement. The search adhered to the Preferred Reporting Items for Systematic Review and Meta-Analyses (PRISMA) guidelines.<sup>14</sup>

### Assessment of quality and heterogeneity

Because of the diversity of methodologies and study designs used, a conventional meta-analysis was not feasible. Therefore, to ensure consistency in the data collected from all the studies included, two authors (Y. K. and M.G.) carefully verified that all variables were reported consistently across all studies, with any differences resolved by the senior author (K.W.). To assess potential bias in the selected studies, we used the National Institutes of Health (NIH) Study Quality Assessment Tools.<sup>15</sup> These tools were specific to certain study designs, and in our review, we used the tools for cohort, case–control, and case series/reports. Two authors (Y. K. and M.G.) independently assessed each study and rated its



**Figure.** Article selection process. Diagram depicts the selection process for studies in this systematic review including databases used and exclusion criteria.

quality as good, fair, or poor based on the assessment tool guidelines. Any discrepancies were resolved through discussion until a consensus was reached.

### Data extraction and statistical analysis

The following data were extracted from each selected study: (1) authors and year of publication, (2) number of patients, (3) age and sex, (4) traumatic/atraumatic causes, (5) mechanism of injury, (6) duration of symptoms before presentation, (7) conservative management, (8) surgical intervention, (9) recovery outcome, and (10) follow-up time. Additionally, all cases were categorized into groups based on (1) cause (traumatic or atraumatic), (2) duration of symptoms before presentation (<1 day, 1–7 days, and 8–84 days), (3) intervention (conservative, surgical, and conservative then surgical), and (4) recovery (full recovery and non-full recovery). Traumatic causes were defined as any injury to the wrist which altered local structure/pressure originating from external forces. Atraumatic causes included any other etiologies that were not a direct result of external forces. Conservative efforts were defined as nonsurgical measures taken to relieve symptoms, most commonly including nonsteroidal anti-inflammatories, arm elevation, steroid injections, and wrist splinting. Other conservative options were targeted toward treating the underlying causes, such as medications for hyperthyroidism, vitamin B6 for deficiency, and diuretics for fluid accumulation. Surgical treatment included carpal tunnel release. Recovery outcome was assessed after intervention as either

**Table 1**  
Traumatic and Atraumatic Causes of ACTS

Traumatic causes	Colle's fracture, burns, ulnar non-union, minor trauma on anticoagulants, injury caused by wrist motion (hyperextension, twisting), MRI-related electrical burn, carpal dislocation, snake bites, cat bites, stonefish envenomation, hemophilia A with trauma, iatrogenic, penetrating wrist trauma, blunt wrist trauma
Atraumatic causes	Infection, spontaneous hemorrhage on anticoagulation, peritendinitis calcarea, autoimmune diseases (rheumatoid arthritis), hyperthyroidism, tophaceous gout, pseudogout, post–upper respiratory tract infection, Rubella immunization, decompression illness, aneurysm of the superficial palmar arch, thrombosis persistent median artery, tumescent fluid with lipodystrophy, post-transplantation, secondary to medications, idiopathic, periarticular calcifications, atypical hand muscles, diabetic myonecrosis, myxofibrosarcoma, malignancy

full resolution or non-full resolution of the symptoms associated with carpal tunnel syndrome.

Pearson's  $\chi^2$  tests were conducted to assess interactions between (1) symptom duration and cause and (2) intervention and cause. A series of logistic regression analyses were performed to investigate and quantify the relationships between (1) recovery outcome and cause, (2) recovery outcome and symptom duration, (3) recovery outcome and intervention, (4) recovery outcome and follow-up duration, and (5) intervention and cause. Additionally, correlation analyses were conducted to assess the associations between (1) follow-up time and cause and (2) follow-up time and intervention. A  $P < .05$  was considered statistical significance for all analyses.

## Results

### Literature search

The literature search yielded 75 articles from PubMed, 295 from Scopus, and 12 from CINAHL. Following the article screen and review, 86 articles were selected for final inclusion. Reasons for exclusion are reported in the PRISMA flowchart (Fig.). The 86 selected articles included 69 case reports, 13 case series, two retrospective, and two prospective studies.<sup>2–4,7,13,16–95</sup>

### Methodological quality of selected articles

For quality assessment, the 86 articles were divided into three groups based on the study design categories established within the NIH Study Quality Assessment Tools: a cohort study (Supplemental Table 1, available on the Journal's website at [www.jhsgo.org](http://www.jhsgo.org)), case–control study (Supplemental Table 2, available on the Journal's website at [www.jhsgo.org](http://www.jhsgo.org)), and a case report/series (Supplemental Table 3 (available on the Journal's website at [www.jhsgo.org](http://www.jhsgo.org)). The studies were graded as follows: 1 for poor quality, 83 for fair quality, and 2 for good quality.

### Characteristics of the collected cases

A total of 197 patients were extracted from collected articles in the age group of 18–91 years, including both males and females of various races. Supplemental Table 4 (available on the Journal's website at [www.jhsgo.org](http://www.jhsgo.org)) summarizes patient demographics, causes of ACTS, duration of symptoms, interventions, follow-up time, and recovery outcomes. Reported causes of ACTS are listed in Table 1 based on traumatic or atraumatic nature. Overall, traumatic causes accounted for approximately 65% of the studied cases. The majority of the cases had symptom duration of 1–7 days (68%), followed by 8–84 days (15%) and <1 day (17%). No significant association was found between cause and symptom duration ( $P = .14$ ). Forty-seven percent were managed conservatively followed by surgery, 30% conservative only, and 23% surgery only. Conservative management followed by surgery was most commonly used in the traumatic group ( $P < .001$ ) but least in the atraumatic group ( $P < .0001$ ). Full recovery was achieved in 168 patients (85%) at

follow-up 91.3 [152.1] days and 29 patients (15%) had a non-full recovery at 121.7 [220.5] days. A summary of the case breakdown based on causes (traumatic or atraumatic) is depicted in Table 2.

### Recovery outcome comparison between groups

Associations between the recovery outcome and follow-up time, duration of symptoms, and intervention were investigated as shown in Table 3. In terms of causes, the atraumatic group was less likely to recover fully compared with the traumatic group (OR: 0.38,  $P = .02$ , 95% CI: 0.17, 0.86) (Table 2), and this association was not affected by the choice of intervention ( $P = .75$ ). Overall, no statistically significant associations were identified between any of the variables: outcome, symptom duration, intervention, and follow-up time (Table 3). When analyzed separately based on causes (atraumatic or traumatic), no significant associations were found between outcome and symptom duration (traumatic:  $P = .41$ , atraumatic:  $P = .31$ ), or outcome and intervention (traumatic:  $P = .23$ , atraumatic:  $P = .38$ ) in either group (Table 4). Full recovery was found to be associated with less follow-up period in the traumatic group ( $P = .04$ ), whereas no significant association was identified between recovery outcome and follow-up duration in the atraumatic group ( $P = .76$ ).

## Discussion

It is not known if traumatic ACTS warrants early surgical decompression or responds to conservative measures and surgeons approach this disease variably, basing it largely on their clinical interpretation and experience. Additionally, many other disease processes can mimic ACTS and further complicate the work-up. Some examples include proximal sites of compression (thoracic outlet syndrome, pronator syndrome), cervical spine diseases, systemic neurological disorders, and nerve contusion injuries.<sup>62,96</sup> Although ACTS can result from a wide variety of etiologies, many advocate that early recognition and prompt surgical intervention, with carpal tunnel release (CTR), are keys to preventing neurovascular damage and optimizing recovery outcomes.<sup>37,62</sup> Although our study did not identify a significant relationship between recovery outcome and symptom durations, we defined symptom duration as the length of time between the onset of symptoms and the patient's presentation to a health care setting. The time it took for patients to be treated after they presented to a provider was not considered. Additionally, symptom duration is subject to variations due to different methods/standards of extraction from each study. Furthermore, because our data do not contain information on treatment duration nor a direct comparison of outcomes stratified by disease severity, we do not have sufficient evidence to conclude associations between recovery outcomes and duration of symptoms. Interestingly, a study performed by Chauhan et al comparing long-term outcomes between elective CTR and acute CTR in combination with open reduction internal fixation of distal radius fractures found no differences between the two cohorts in symptom severity, functional outcomes, and Boston Carpal Tunnel Questionnaire scores.<sup>97</sup> However, the degree of improvement was

**Table 2**  
Case Breakdown per Cause

Variables		Traumatic ACTS (n = 127, 64.5%)	Atraumatic ACTS (n = 70, 35.5%)	P Values
Symptom duration (n, % total)	<1 d	10 (15.2%)	13 (18.6%)	0.14
	1–7 d	50 (75.8%)	43 (61.4%)	
	8–84 d	6 (9.1%)	14 (20%)	
Intervention (n, % total)	Conservative only	33 (26%)	23 (32.9%)	<b>0.003*</b>
	Surgical only	22 (17.3%)	24 (34.3%)	
	Conservative then surgical	72 (56.7%)	23 (32.9%)	
Recovery (n, % total)	Full	114 (89.8%)	54 (77.1%)	<b>0.02*</b>
	Non-full	13 (10.2%)	16 (22.9%)	
	Follow-up (d)	91 (45.6 – 182.5)	114 (37.7 – 205.3)	

\* Represents statistical significance.

**Table 3**  
Associations between Variables

Variables	P Values
Outcome and symptom duration	.83
Outcome and intervention	.17
Outcome and follow-up time	.26
Follow-up time and symptom duration	.52
Follow-up time and intervention	.29

**Table 4**  
Associations between Outcome and Symptom Duration, Intervention, and Follow-Up Time Separated by Cause

Subgroup	Variables	P Values
Traumatic	Outcome and symptom duration	.41
	Outcome and intervention	.23
	Outcome and follow-up time	<b>.04*</b>
Atraumatic	Outcome and symptom duration	.31
	Outcome and intervention	.38
	Outcome and follow-up time	.76

\* Represents statistical significance.

not determined between the two cohorts due to the lack of pre-operative assessment scores. Therefore, it is possible that severity also affects clinical outcomes, although this specific determination is yet to be delineated.

Ford and Ali reported that patients recovered fully if surgical interventions were performed promptly after symptom onset.<sup>37</sup> Mack et al recommended approaching suspected ACTS first with conservative measures such as elevation and splinting for 2 hours.<sup>62</sup> If symptoms progress, carpal tunnel pressure should be measured, and CTR should be performed within 8 hours of onset if the intracarpal tunnel pressure exceeds 40 mmHg. However, although thought to be the treatment of choice by many, there is no universal consensus on the timeframe for which CTR should be performed.<sup>98</sup> Furthermore, it is important to note that Ford and Ali,<sup>37</sup> and Mack et al<sup>62</sup> only included ACTS resulting from traumatic fractures of the distal wrist/hand. As the characteristics of traumatic and atraumatic injuries are innately different, this raises the question of whether or not the conclusions made by the two studies apply to atraumatic ACTS. Although recommending operative decompression overall, Adamson et al indicated that conservative treatments may be considered in treating atraumatic and non-severe traumatic ACTS.<sup>16</sup> Our analysis indicated that although patients with traumatic etiologies received better recovery overall, the outcomes were not dependent on the choice of intervention in either traumatic or atraumatic groups.

In atraumatic cases, the precipitating medical conditions play a large role in recovery outcomes, depending on their severity, and treatments should target both managing medical conditions and

providing symptomatic relief. Padua et al successfully treated bilateral ACTS with methimazole and propranolol in a patient with undiagnosed hyperthyroidism.<sup>72</sup> The patient experienced complete resolution of symptoms at 7 months of follow-up. Mayne et al<sup>65</sup> and Rahmtoola et al<sup>78</sup> reported treating ACTS secondary to wrist trauma in hemophilia patients with factor VIII followed by CTR, resulting in complete recovery in all cases. However, with infectious etiologies, although atraumatic, management options vary depending on the clinical course and source of infection. Sahs et al<sup>99</sup> mainly treated ACTS secondary to toxic shock syndrome conservatively, whereas Yoshida et al<sup>94</sup> and Wilhelm et al<sup>95</sup> treated a localized infection such as tenosynovitis operatively. Other examples of conservative approaches in treating atraumatic cases included steroid injections, removal of inciting medications, rest, splinting, and hyperbaric oxygen in decompression illness.<sup>13,33,46,58</sup> However, there is a lack of evidence demonstrating the optimal timeframe for conservative management. Our results indicate that atraumatic ACTS is less likely to recover fully compared with traumatic etiologies. A likely explanation could be that the slower onset of symptoms that accompanies many atraumatic cases may not prompt patients to seek care immediately, which in turn results in more permanent damage to the nerve. Conversely, traumatic causes typically result in more obvious structural changes in a shorter period and may alert patients to seek care promptly. Although the difference was not statistically significant, we observed that a greater proportion of patients in the atraumatic group (20%) had symptoms lasting more than 8 days, compared with 9.1% in the traumatic cohort.

In terms of recovery time, patients have reported relief of ACTS symptoms as early as 1–5 days after intervention, regardless of intervention or whether the etiology was traumatic or atraumatic.<sup>16,21,60,66,86</sup> Conversely, some patients have experienced persistent symptoms for more than 1 year after intervention, independent of the underlying causes of the disease.<sup>23,56,66,99</sup> In our series, we found no significant differences in follow-up duration between the traumatic and atraumatic subgroups, suggesting that the more favorable outcomes observed in the traumatic cohort were because of the traumatic nature rather than the time effect. Overall, there was a trend of longer follow-up duration in those without a full recovery (121.7 [220.5] days vs 91.3 [152.1] days), and this difference became significant within the traumatic subgroup. Although follow-up duration does not necessarily correlate the recovery time, incorporating this variable into statistical analyses helps mitigate the potential confounding effects caused by the time factor.

This review is limited by the types of articles included in the final analysis. An overwhelming majority of the articles are case reports and case series, which provide lower levels of evidence compared with other types of clinical studies. Especially, there are no randomized controlled trials or comparative studies to assess and compare ACTS treatments. Additionally, although the sample

size of 197 was sufficient for logistic regression analyses with only one predictor,<sup>100</sup> it may not be adequate for logistic regression analyses with an interaction term. Many of the analyzed articles were published more than 10 years ago, highlighting the likely outdated information and questioning the validity of the treatment approaches reported in each study. Notably, it is possible that some patients included in the study had median neuropathy instead of ACTS, especially in traumatic cases. Patients with median neuropathy present similarly to ACTS patients but with a lesser degree of nerve damage, thus the management and prognosis are different than those with ACTS.<sup>100</sup> The determination of final recovery status was based on the reported data from each study, which may have slight variations due to differences in assessment methods and author interpretation. Finally, although all ACTS cases were considered as a single disease, the severity of each case was not considered, which could potentially have effects on recovery.

Traumatic ACTS is associated with better recovery outcomes than atraumatic causes. Prompt surgical intervention does not result in better recovery compared with a conservative approach, regardless of traumatic or atraumatic etiologies. As this study was limited by the predominance of case reports and case studies, further investigation with prospective studies is required to determine whether recovery outcomes differ between surgical and conservative management in the treatment of ACTS.

## References

- Aroori S, Spence RA. Carpal tunnel syndrome. *Ulster Med J*. 2008;77(1):6–17.
- Fricker R, Fuhr P, Pippert H, Troeger H. Acute median nerve compression at the distal forearm caused by a thrombosed aneurysm of an epineural vessel: case report. *Neurosurgery*. 1996;38(1):194–196.
- Martin SD, Sharrock NE, Mineo R, Sobel M, Weiland AJ. Acute exacerbation of carpal tunnel syndrome after radial artery cannulation. *J Hand Surg Am*. 1993;18(3):455–458.
- Weiber H, Linell F. Tumoral calcinosis causing acute carpal tunnel syndrome. *Case report. Scand J Plast Reconstr Surg Hand Surg*. 1987;21(2):229–230.
- Jhattu H, Klaassen S, Ying C, Hussain MA. Acute carpal tunnel syndrome in trauma. *Eur J Plast Surg*. 2012;35(9):639–646.
- Schnetzler KA. Acute carpal tunnel syndrome. *JAAOS - J Am Acad Orthop Surg*. 2008;16(5):276–282.
- Leow JM, Clement ND, McQueen MM, Duckworth AD. The rate and associated risk factors for acute carpal tunnel syndrome complicating a fracture of the distal radius. *Eur J Orthop Surg Traumatol*. 2021;31(5):981–987.
- Tosti R, Ilyas AM. Acute carpal tunnel syndrome. *Orthop Clin North Am*. 2012;43(4):459–465.
- Szabo RM. Acute carpal tunnel syndrome. *Hand Clin*. 1998;14(3):419–429.
- Atroshi I, Gummesson C, Johnsson R, Ornstein E, Ranstam J, Rosén I. Prevalence of carpal tunnel syndrome in a general population. *JAMA*. 1999;282(2):153–158.
- Pourmemari MH, Heliövaara M, Viikari-Juntura E, Shiri R. Carpal tunnel release: lifetime prevalence, annual incidence, and risk factors. *Muscle Nerve*. 2018;58(4):497–502. <https://doi.org/10.1002/mus.26145>
- Michelsen H, Posner MA. Medical history of carpal tunnel syndrome. *Hand Clin*. 2002;18(2):257–268.
- Hale MS, Ruderman JE. Carpal tunnel syndrome associated with rubella immunization. *Am J Phys Med*. 1973;52(4):189–194.
- Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *BMJ*. 2009;339:b2535.
- Study Quality Assessment Tools | NHLBI, NIH. Accessed May 7, 2023. <https://www.nlm.nih.gov/health-topics/study-quality-assessment-tools>
- Adamson JE, Srouji SJ, Horton CE, Mladick RA. The acute carpal tunnel syndrome. *Plast Reconstr Surg*. 1971;47(4):332–336.
- Ahmad F, Michalski J, Winterton M, Simcock X, Wysocki RW. Spontaneous diabetic myonecrosis presenting as acute carpal tunnel syndrome. *J Hand Surg Glob Online*. 2022;4(1):53–56.
- Ametewee K, Harris A, Samuel M. Acute carpal tunnel syndrome produced by anomalous flexor digitorum superficialis indicis muscle. *J Hand Surg Br*. 1985;10(1):83–84.
- Aslam N, Lo S, McNab I. Acute carpal tunnel syndrome caused by a fragment of bone from metacarpal injury. *Inj Extra*. 2005;36(4):73–76.
- Bache SE, Taggart I. Acute carpal tunnel syndrome following a burn in a patient with McArdle's disease: a case report. *Burns*. 2012;38(7):e17–e19.
- Balakrishnan C, Smith MF, Puri P. Acute carpal tunnel syndrome from thrombosed persistent median artery. *J Emerg Med*. 1999;17(3):437–439.
- Barbee GA, Haley CL, Berry-Cabán CS. A case of acute carpal tunnel syndrome. *JAAPA*. 2016;20(1):31–32.
- Bauman T, Gelbermann R, Mubarak S, Garfin S. The acute carpal tunnel syndrome. *Clin Orthop Relat Res*. 1981;6(156):151–156.
- Belsole RJ, Greeley JM. Surgeon's acute carpal tunnel syndrome: an occupational hazard? *J Fla Med Assoc*. 1988;75(6):369–370.
- Black PRM, Flowers MJ, Saleh M. Acute carpal tunnel syndrome as a complication of oral anticoagulant therapy. *J Hand Surg*. 1997;22(1):50–51.
- Bonatz E, Seabolt KE. Acute carpal tunnel syndrome in a patient taking Coumadin: case report. *J Trauma*. 1993;35(1):143–144.
- Boström L, Svartengren G. Acute carpal tunnel syndrome caused by peritendinitis calcarea: two case reports. *Scand J Plast Reconstr Surg Hand Surg*. 1993;27(2):157–159.
- Charalambous CP, Zipitis CS, Kumar R, Paul AS. Acute carpal tunnel syndrome: two rare cases. *Hand Surg*. 2003;08(1):117–118.
- Chiu KY, Ng WF, Wong WB, Choi CH, Chow SP. Acute carpal tunnel syndrome caused by pseudogout. *J Hand Surg Am*. 1992;17(2):299–302.
- Copeland J, Wells HG, Puckett CL. Acute carpal tunnel syndrome in a patient taking coumadin. *J Trauma*. 1989;29(1):131–132.
- Dahmam A, Matter-Parrat V, Manguila F, Giannikas D, Braun FM. Acute carpal tunnel syndrome due to a thrombosed persistent median artery: unusual cause in athletes. *J Traumatol Sport*. 2015;32(3):126–128.
- Din R, Giannikas K, El-Hadidi M. Acute calcifying tendonitis – an unusual cause of carpal tunnel syndrome. *Eur J Emerg Med*. 2001;8(1):65–66.
- Eisenbud L, Ejadi S, Mar N. Development of carpal tunnel syndrome in association with checkpoint inhibitors. *J Oncol Pharm Pract*. 2021;27(3):764–765.
- El Hajj II, Harb MI, Sawaya RA. Acute progressive bilateral carpal tunnel syndrome after upper respiratory tract infection. *South Med J*. 2005;98(11):1149–1151.
- Faithfull DK, Wallace RF. Traumatic rupture of median artery an unusual cause for acute median nerve compression. *J Hand Surg Br*. 1987;12(2):233–235.
- Figus A, Iwuagwu FC, Elliot D. Subacute nerve compressions after trauma and surgery of the hand. *Plast Reconstr Surg*. 2007;120(3):705–712.
- Ford DJ, Ali MS. Acute carpal tunnel syndrome. Complications of delayed decompression. *J Bone Joint Surg Br*. 1986;68(5):758–759.
- Gallagher B, Khalifa M, Van Heerden P, Elbardsy N. Acute carpal tunnel syndrome due to filarial infection. *Pathol Res Pract*. 2002;198(1):65–67.
- Garriguez-Pérez D, Serrano-Mateo L, Donadeu-Sánchez S, Gimeno MD, Marco F. Acute carpal tunnel syndrome after radial artery cannulation: a case report. *JBJS Case Connect*. 2022;12(1):e21.00495.
- Gaur SC, Kulshreshtha K, Swarup S. Acute carpal tunnel syndrome in Hansen's disease. *J Hand Surg Br*. 1994;19(3):286–287.
- Gerardi JA, Mack GR, Lutz RB. Acute carpal tunnel syndrome secondary to septic arthritis of the wrist. *J Am Osteopath Assoc*. 1989;89(7):933–934.
- Guyon MA, Honet JC. Carpal tunnel syndrome or trigger finger associated with neck injury in automobile accidents. *Arch Phys Med Rehabil*. 1977;58(7):325–327.
- Hargreaves DG, Gosal H, Moss AL. Another cause for acute carpal tunnel syndrome: tricyclic overdose. *J Accid Emerg Med*. 1995;12(2):158–159.
- Head LK, Bradley R, Momtazi M. Tenosynovial chondromatosis of the wrist presenting with acute carpal tunnel syndrome: a case report. *Hand Surg Rehabil*. 2018;37(2):117–120.
- Howie CR, Buxton R. Acute carpal tunnel syndrome due to spontaneous haemorrhage. *J Hand Surg Br*. 1984;9(2):137–138.
- Isakov AP, Broome JR, Dutka AJ. Acute carpal tunnel syndrome in a diver: evidence of peripheral nervous system involvement in decompression illness. *Ann Emerg Med*. 1996;28(1):90–93.
- Jacob ZC, Tito MF, Dagum AB. MR imaging-related electrical thermal injury complicated by acute carpal tunnel and compartment syndrome: case report. *Radiology*. 2010;254(3):846–850.
- Jones WA, Schecher LR. Acute carpal tunnel syndrome: a case report. *J Hand Surg Br*. 1988;13(4):400–401.
- Kannan A, Khatri D, Trikha V, Choudhary B. Isolated volar fracture-dislocation of the scaphoid with acute carpal tunnel syndrome: a case report. 2010;76(4):552–554.
- Khashaba A. Carpal tunnel syndrome from thrombosed persistent median artery. *J Emerg Med*. 2002;22(1):55–57.
- Knight DJ, Gibson PH. Acute calcification and carpal tunnel syndrome. *J Hand Surg Br*. 1993;18(3):335–336.
- Knobloch K, Gohritz A, Altintas MA, Spies M, Vogt PM. A wakeboarding injury presented as acute carpal syndrome and median nerve contusion after wrist strangulation: a case report. *Cases J*. 2009;2(1):100.
- Kono H. Acute carpal tunnel syndrome caused by anomalous muscle bellies: a case report. *Hand Surg*. 2003;08(1):141–143.
- Lambe G, Scott S, Acharya A. Closed rupture of a lumbrical muscle. *Eur J Plast Surg*. 2002;25(2):97–98.
- Larson BJ, DeLange LC. Traumatic volar dislocation of the trapezoid with acute carpal tunnel syndrome. *Orthopedics*. 2005;28(2):165–167.
- Lazaro RP. Complex regional pain syndrome and acute carpal tunnel syndrome following radial artery cannulation: a neurological perspective and review of the literature. *Medicine*. 2015;94(3):e422.
- Lee L, Yao J. Stenosing flexor tenosynovitis following a rattlesnake bite. *Orthopedics*. 2010;33(7):515.
- Lewis RA, Shea OF, Shea KG. Acute carpal tunnel syndrome: wrist stress during a major climb. *Phys Sportsmed*. 1993;21(7):102–108.

59. Ling SKK, Cheng SC, Yen CH. Stonefish envenomation with acute carpal tunnel syndrome. *Hong Kong Med J*. 2009;15(6):471–473.
60. Lombardi AS, Quirke TE, Rauscher G. Acute median nerve compression associated with tumescent fluid administration. *Plast Reconstr Surg*. 1998;102(1):235–237.
61. Lourie GM, Scott Levin L, Toby B, Urbaniak J. Distal rupture of the palmaris longus tendon and fascia as a cause of acute carpal tunnel syndrome. *J Hand Surg*. 1990;15(2):367–369.
62. Mack GR, McPherson SA, Lutz RB. Acute median neuropathy after wrist trauma. The role of emergent carpal tunnel release. *Clin Orthop Relat Res*. 1994;300:141–146.
63. Mahmud T. Bilateral acute carpal tunnel syndrome after combined pancreatic and renal transplant. *Scand J Plast Reconstr Surg Hand Surg*. 2009;43(3):174–176.
64. Maxwell JA, Ketchum LD. Acute carpal tunnel syndrome secondary to thrombosis of a persistent median artery. *J Neurosurg*. 1973;38:4.
65. Mayne AIW, Howard A, Kent M, Banks J. Acute carpal tunnel syndrome in a patient with haemophilia. *BMJ Case Rep*. 2012;2012:bcr0320126152.
66. McClain EJ, Wissinger HA. The acute carpal tunnel syndrome: nine case reports. *J Trauma*. 1976;16(1):75–78.
67. Nather A, Chacha PB, Lim P. Acute carpal tunnel syndrome secondary to thrombosis of a persistent median artery (with high division of the median nerve). A case report. *Ann Acad Med Singap*. 1980;9(1):118–121.
68. Nettrour JF, Eggers SD, Pittelkow MR, Matteson EL. Acute carpal tunnel syndrome preceded by 5 years of unusual skin changes. *Arch Neurol*. 2007;64(3):447.
69. Nkele C. Acute carpal tunnel syndrome resulting from haemorrhage into the carpal tunnel in a patient on warfarin. *J Hand Surg Br*. 1986;11(3):455–456.
70. Nourissat G, Fournier E, Werther JR, Dumontier C, Doursounian L. Acute carpal tunnel syndrome secondary to pyogenic tenosynovitis. *J Hand Surg*. 2006;31(6):687–688.
71. Olerud C, Lönnquist L. Acute carpal tunnel syndrome caused by fracture of the scaphoid and the 5th metacarpal bones. *Injury*. 1984;16(3):198–199.
72. Padua L, Monaco M, Gregori B, Tonalì PA. Bilateral acute carpal tunnel syndrome and hyperthyroidism: a case report. *Eur J Neurol*. 1996;3(4):399–401.
73. Pai CH, Howard Tseng C. Acute carpal tunnel syndrome caused by tophaceous gout. *J Hand Surg*. 1993;18(4):667–669.
74. Pai V, Pai V, Muir R. Periarticular calcification causing acute carpal tunnel syndrome: a case report. *J Orthop Surg (Hong Kong)*. 2009;17(2):234–237.
75. Pai V. Acute carpal tunnel syndrome complicating infective tenosynovitis at the wrist: a case report. *J Orthop Surg*. 1996;4(1):95–98.
76. Parthenis DG, Karagkevrekis CB, Waldram MA. Von Willebrand's disease presenting as acute carpal tunnel syndrome. *J Hand Surg*. 1998;23(1):114.
77. Piereschi S, Remy H, Camuzard O. Volar fracture dislocation of the 2nd metacarpal base associated with acute carpal tunnel syndrome: a case report. *Ann Chir Plast Esthét*. 2021;66(2):180–183.
78. Rahimtoola ZO, van Baal SG. Two cases of acute carpal tunnel syndrome in classic haemophilia. *Scand J Plast Reconstr Surg Hand Surg*. 2002;36(3):186–188.
79. Rate AJ, Parkinson RW, Meadows TH, Freemont AJ. Acute carpal tunnel syndrome due to pseudogout. *J Hand Surg Br*. 1992;17(2):217–218.
80. Reddy KJ, Packer GJ. Stabilization of an acute perilunate dislocation using the “tag” suture anchor. *J Hand Surg*. 1998;23(2):262–263.
81. Rose RE. Acute carpal tunnel syndrome secondary to thrombosis of a persistent median artery. *West Indian Med J*. 1995;44(1):32–33.
82. Ross C. Vitamin B6 and carpal tunnel syndrome a case report. *J Orthomol Med*. 1994;9(1):49–50.
83. Saleh WR, Yajima H, Nakanishi A. Acute carpal tunnel syndrome secondary to calcific tendinitis: case report. *Hand Surg*. 2008;13(03):197–200.
84. Samii K, Cassinotti P, de Freudenreich J, Gallopin Y, Le Fort D, Stalder H. Acute bilateral carpal tunnel syndrome associated with human parvovirus B19 infection. *Clin Infect Dis*. 1996;22(1):162–164.
85. Sbai MA, Dabloun S, Benzarti S, Khechimi M, Jenzeri A, Maalla R. Acute carpal tunnel syndrome of the hand following a cat bite. *Pan Afr Med J*. 2015;21:206.
86. Seiler JG, Havig M, Carpenter W. Acute carpal tunnel syndrome complicating chronic palmar subluxation of the distal ulna. *J South Orthop Assoc*. 1996;5(2):108–110.
87. Sibley PA, Mandel RJ. Atraumatic acute carpal tunnel syndrome in a patient taking dabigatran. *Orthopedics*. 2012;35(8):e1286–e1289.
88. Simon DA, Taylor TL. Horseshoe abscess associated with acute carpal tunnel syndrome: somebody wake up the hand surgeon. *CJEM*. 2012;14(2):124–127.
89. Slesarenko Y, Dagum AB, Hurst L. False aneurysm of the superficial palmar arch causing acute carpal tunnel syndrome. *Orthopedics*. 2007;30(6):493–494.
90. Sterling AP, Eshraghi A, Anderson WJ, Habermann ET. Acute carpal tunnel syndrome secondary to a foreign body within the median nerve. *Bull Hosp Joint Dis*. 1972;33(2):130–134.
91. Unglaub F, Wolf MB, Dragu A, Horch RE. Bilateral atypical muscles causing acute bilateral carpal tunnel syndrome in recreational climber. *Arch Orthop Trauma Surg*. 2010;130(1):37–40.
92. Verfaillie stefaan, De Smet L, Leemans A, Van Damme B, Fabry G. Acute carpal tunnel syndrome caused by hydroxyapatite crystals: a case report. *J Hand Surg*. 1996;21(3):360–362.
93. Wadström J, Gannedahl G, Claesson K, Wahlberg J. Acute carpal tunnel syndrome immediately after combined kidney and pancreas transplantation. *Transplant Proc*. 1995;27(6):3489–3490.
94. Wilhelm A, Romeo N, Trevino R. A rare cause of pyogenic flexor tenosynovitis: nocardia nova. *J Hand Surg*. 2018;43(8):778.e1–778.e4.
95. Yoshida H, Imura H, Goto T, Nakamata T, Daya MR, Kamiya T. Acute carpal tunnel syndrome due to pyogenic flexor tenosynovitis without any antecedent injury. *Intern Med*. 2017;56(11):1439–1442.
96. Dengler J, Stephens JD, Bamberger HB, Moore AM. Mimickers of carpal tunnel syndrome. *JBJS Rev*. 2020;8(2):e0087.
97. Chauhan A, Bowlin TC, Mih AD, Merrell GA. Patient-reported outcomes after acute carpal tunnel release in patients with distal radius open reduction internal fixation. *Hand (N Y)*. 2012;7(2):147–150.
98. Gillig JD, White SD, Rachel JN. Acute carpal tunnel syndrome: a review of current literature. *Orthop Clin North Am*. 2016;47(3):599–607.
99. Sahs AL, Helms CM, DuBois C. Carpal tunnel syndrome. Complication of toxic shock syndrome. *Arch Neurol*. 1983;40(7):414–415.
100. Holbrook HS, Hillesheim RA, Weller WJ. Acute carpal tunnel syndrome and median nerve neuropathia: a review. *Orthop Clin North Am*. 2022;53(2):197–203.