

CLINICAL PRACTICE GUIDELINE ON THE TREATMENT OF CARPAL TUNNEL SYNDROME

Adopted by the American Academy of Orthopaedic Surgeons

Board of Directors

September 2008

This clinical guideline was developed by an AAOS physician volunteer Work Group and is provided as an educational tool based on an assessment of the current scientific and clinical information and accepted approaches to treatment. It is not intended to be a fixed protocol as some patients may require more or less treatment. Patient care and treatment should always be based on a clinician's independent medical judgment given the individual clinical circumstances.

Endorsed By:











DISCLAIMER

This Clinical Practice Guideline was developed by an AAOS physician volunteer Work Group based on a systematic review of the current scientific and clinical information and accepted approaches to treatment and/or diagnosis. This Clinical Practice Guideline is not intended to be a fixed protocol, as some patients may require more or less treatment or different means of diagnosis. Clinical patients may not necessarily be the same as those found in a clinical trial. Patient care and treatment should always be based on a clinician's independent medical judgment, given the individual patient's clinical circumstances.

CONFLICT OF INTEREST

All panel members gave full disclosure of conflicts of interest prior to voting on the recommendations contained within these guidelines.

FUNDING SOURCE

These guidelines were funded exclusively by the American Academy of Orthopaedic Surgeons who received no funding from outside commercial sources to support the development of this document.

AAOS Clinical Practice Guideline Copyright

Disclaimer

This Clinical Practice Guideline was developed by an AAOS physician volunteer Work Group based on a systematic review of the current scientific and clinical information and accepted approaches to treatment and/or diagnosis. This Clinical Practice Guideline is not intended to be a fixed protocol, as some patients may require more or less treatment or different means of diagnosis. Clinical patients may not necessarily be the same as those found in a clinical trial. Patient care and treatment should always be based on a clinician's independent medical judgment, given the individual patient's clinical circumstances.

Disclosure Requirement

In accordance with AAOS policy, all individuals whose names appear as authors or contributors to Clinical Practice Guideline filed a disclosure statement as part of the submission process. All panel members provided full disclosure of potential conflicts of interest prior to voting on the recommendations contained within this Clinical Practice Guidelines.

Funding Source

This Clinical Practice Guideline was funded exclusively by the American Academy of Orthopaedic Surgeons who received no funding from outside commercial sources to support the development of this statement.

FDA Clearance

Some drugs or medical devices referenced or described in this Clinical Practice Guideline may not have been cleared by the Food and Drug Administration (FDA) or may have been cleared for a specific use only. The FDA has stated that it is the responsibility of the physician to determine the FDA clearance status of each drug or device he or she wishes to use in clinical practice.

Copyright

All rights reserved. No part of this Clinical Practice Guideline may be reproduced, stored in a retrieval system, or transmitted, in any form, or by any means, electronic, mechanical, photocopying, recording, or otherwise, without prior written permission from the AAOS.

Published 2008 by the American Academy of Orthopaedic Surgeons 6300 North River Road Rosemont, IL 60018 First Edition Copyright 2008 by the American Academy of Orthopaedic Surgeons

TABLE OF CONTENTS

SUMMARY OF RECOMMENDATIONSV					
WORK GROUP PANELVIII					
I. OVERVIEW1					
Goals and Rationale					
Scope and Organization2					
Intended Users2					
Patient Population					
Incidence and Prevalence					
Burden of Disease					
Etiology					
Diagnosis and treatment					
Risk Factors					
II. METHODS4					
Process Overview4					
Consensus Development					
Article Inclusion and Exclusion Criteria					
Literature Searches					
Assigning a Level of Evidence					
Data Extraction					
Grading the Recommendations					
Statistical Methods					
Revision Plans					

III.	RECOMMENDATIONS	10
Recon	nmendation 1	. 10
Recon	nmendation 2	. 11
Recon	nmendation 3	. 12
Recon	nmendation 4a	. 13
Recon	nmendation 4b	. 13
Recon	nmendation 4c	. 14
Recon	nmendation 4d	. 15
Recon	nmendation 4e	. 15
Recon	nmendation 5	. 17
Recon	nmendation 6	. 18
Recon	nmendation 7	. 19
Recon	nmendation 8	. 20
Recon	nmendation 9	. 21
Futur	e Research	. 23
Refer	ences	. 24
Refer	ences to Included Articles	. 25
Refer	ences to Excluded Articles	. 33
IV.	APPENDIXES	53
Appei	ndix I: Literature Searches	. 54
Appei	ndix II: Article Inclusions and Exclusions	. 59
Appei	ndix III: Rating Evidence Quality	. 70
Appei	ndix IV: Evidence Tables	. 71
Appei	ndix V: Conflicts of Interest	. 74
Appei	ndix VI: Documentation of Approval	. 75
Appei	ndix VII: Advisory Review Panel	. 76

Summary of Recommendations

The following is a summary of the recommendations in the AAOS' clinical practice guideline, The Treatment of Carpal Tunnel Syndrome. This summary does not contain rationales that explain how and why these recommendations were developed nor does it contain the evidence supporting these recommendations. All readers of this summary are strongly urged to consult the full guideline and evidence report for this information. We are confident that those who read the full guideline and evidence report will also see that the recommendations were developed using systematic evidence-based processes designed to combat bias, enhance transparency, and promote reproducibility. This summary of recommendations is not intended to stand alone.

Recommendation 1

A course of non-operative treatment is an option in patients diagnosed with carpal tunnel syndrome. Early surgery is an option when there is clinical evidence of median nerve denervation or the patient elects to proceed directly to surgical treatment.

(Grade C, Level V)

Recommendation 2

We suggest another non-operative treatment or surgery when the current treatment fails to resolve the symptoms within 2 weeks to 7 weeks.

(Grade B, Level I and II)

Recommendation 3

We do not have sufficient evidence to provide specific treatment recommendations for carpal tunnel syndrome when found in association with the following conditions: diabetes mellitus, coexistent cervical radiculopathy, hypothyroidism, polyneuropathy, pregnancy, rheumatoid arthritis, and carpal tunnel syndrome in the workplace.

(Inconclusive, No evidence found)

Recommendation 4a

Local steroid injection or splinting is suggested when treating patients with carpal tunnel syndrome, before considering surgery.

(Grade B, Level I and II)

Recommendation 4b

Oral steroids or ultrasound are options when treating patients with carpal tunnel syndrome.

(Grade C, Level II)

Recommendation 4c

We recommend carpal tunnel release as treatment for carpal tunnel syndrome.

(Grade A, Level I)

Recommendation 4d

Heat therapy is not among the options that should be used to treat patients with carpal tunnel syndrome.

(Grade C, Level II)

Recommendation 4e

The following treatments carry no recommendation for or against their use: activity modifications, acupuncture, cognitive behavioral therapy, cold laser, diuretics, exercise, electric stimulation, fitness, graston instrument, iontophoresis, laser, stretching, massage therapy, magnet therapy, manipulation, medications (including anticonvulsants, antidepressants and NSAIDs), nutritional supplements, phonophoresis, smoking cessation, systemic steroid injection, therapeutic touch, vitamin B6 (pyridoxine), weight reduction, yoga.

(Inconclusive, Level II and V)

Recommendation 5

We recommend surgical treatment of carpal tunnel syndrome by complete division of the flexor retinaculum regardless of the specific surgical technique.

(Grade A, Level I and II)

Recommendation 6

We suggest that surgeons do not routinely use the following procedures when performing carpal tunnel release:

```
skin nerve preservation (Grade B, Level I) epineurotomy (Grade C, Level II)
```

The following procedures carry no recommendation for or against use: flexor retinaculum lengthening, internal neurolysis, tenosynovectomy, ulnar bursa preservation

(Inconclusive, Level II and V).

Recommendation 7

The physician has the option of prescribing pre-operative antibiotics for carpal tunnel surgery.

(Grade C, Level III)

Recommendation 8

We suggest that the wrist not be immobilized postoperatively after routine carpal tunnel surgery

```
(Grade B, Level II).
```

We make no recommendation for or against the use of postoperative rehabilitation.

(Inconclusive, Level II).

Recommendation 9

We suggest physicians use one or more of the following instruments when assessing patients' responses to CTS treatment for research:

- Boston Carpal Tunnel Questionnaire (disease-specific)
- DASH Disabilities of the arm, shoulder, and hand (region-specific; upper limb)
- MHQ Michigan Hand Outcomes Questionnaire (region-specific; hand/wrist)
- PEM (region-specific; hand)
- SF-12 or SF-36 Short Form Health Survey (generic; physical health component for global health impact)

(Grade B, Level I, II, and III)

Work Group Panel

Michael Warren Keith, MD (Chair) 2500 Metro Health Drive Cleveland, OH 44109-1900 Orthopaedic Hand Surgeon

Victoria Masear, MD (Co-Chair) 48 Medical Park E Dr Ste 255 Birmingham, AL 35235-3411 Orthopaedic Hand Surgeon

Kevin Chung, MD University of Michigan Medical Center 1500 East Medical Center Drive 2130 Taubman Health Care Center Ann Arbor, MI 48109-0340 Plastic and Reconstructive Surgery

Peter C Amadio, MD Mayo Clinic 200 1st St S W Rochester, MN 55902-3008 Orthopaedic Hand Surgeon

Michael Andary, MD Michigan State University B401 W Fee Hall (PMR) East Lansing, MI 48824-1316 Physical Medicine and Rehabilitation Neurology

Richard W. Barth, MD 2021 K St Ste 400 Washington, DC 20006-1003 AAOS Board of Councilors Orthopaedic Hand Surgeon

Kent Maupin, MD 1111 Leffingwell NE Ste 200 Grand Rapids, MI 49525 Orthopaedic Surgery

Brent Graham MD

University of Toronto 399 Bathurst St. 425-2 East Wing Toronto, ON M5T 2S8 Canada Orthopaedic Hand Surgeon/Microsurgery Guidelines Oversight Chair: William C. Watters, III MD 6624 Fannin #2600 Houston, TX 77030 Orthopaedic Spine Surgeon

AAOS Staff: Charles M. Turkelson, PhD AAOS Research Director 6300 N River Road Rosemont, IL 60018

Robert H. Haralson III, MD, MBA AAOS Medical Director 6300 N River Road Rosemont, IL 60018 Orthopaedic Surgeon

Janet L. Wies MPH Clinical Practice Guideline Mgr 6300 N River Road Rosemont, IL 60018

AAOS Research Analysts Kevin Boyer – Lead Analyst Andrew Chang MPH Erica Smith MPH

AAOS Summer Interns Andy Riff Jacqueline Roache MPH

AMERICAN ACADEMY OF ORTHOPAEDIC SURGEONS CLINICAL PRACTICE GUIDELINE ON THE TREATMENT OF CARPAL TUNNEL SYNDROME

I OVERVIEW

Evidence-based Practice (EBP) standards are in a state of continuous evolution. Current EBP standards demand that physicians use the best available evidence to guide their clinical decision making processes. Increasingly rigorous EBP standards have also resulted in more rigorous clinical studies of ever stronger design, complexity, and statistical analysis. This clinical practice guideline consists of a systematic review of the available literature regarding the treatment of carpal tunnel syndrome. The purpose of this clinical practice guideline is to help improve carpal tunnel syndrome treatment based on the current best evidence. The systematic review detailed herein was conducted between June and October of 2007 and demonstrates where there is good evidence, where evidence is lacking, and what topics future research must target in order to improve carpal tunnel syndrome treatment. The AAOS Carpal Tunnel Syndrome (CTS) Guideline Work Group systematically reviewed the available literature, evaluated the level of evidence found in that literature, and subsequently wrote the following recommendations based on a rigorous, standardized process.

GOALS AND RATIONALE

The AAOS has created this clinical practice guideline to improve patient care by outlining the appropriate information-gathering and decision-making processes involved in managing the treatment of carpal tunnel syndrome. This guideline is also an educational tool to guide qualified physicians (see Intended Users) through a series of treatment decisions in an effort to improve the quality and efficiency of care.

This guideline should not be construed as including all proper methods of care or excluding methods of care reasonably directed to obtaining the same results. The ultimate judgment regarding any specific procedure or treatment must be made in light of all circumstances presented by the patient and the needs and resources particular to the locality or institution. Further, the patient must be an active participant in treatment decisions. All treatment for CTS is based on the assumption that final decisions are predicated on patient and physician mutual communication about available treatment alternatives and procedures applicable to the individual patient. These decisions include an evaluation of the patient's current quality of life with CTS. Patients will present with considerable variability in acceptable choices, needs, and access to non-operative alternatives. It is understood that after the patient has been informed of available alternative non-operative therapies and has discussed these options with their physician, the informed patient choice may be to go directly to surgery.

SCOPE AND ORGANIZATION

INTENDED USERS

This guideline is intended to be used by all appropriately trained surgeons and all qualified physicians considering treatment of CTS. Typically, appropriately trained surgeons will have completed medical training, a qualified residency and some may have additional sub-specialty training. Insurance payers, governmental bodies, and health-policy decision-makers may also find this guideline useful as an evolving standard of evidence regarding treatment of Carpal Tunnel Syndrome.

PATIENT POPULATION

Persons of all genders, races, ages, occupations and health status may be afflicted by Carpal Tunnel Syndrome. The present guideline is aimed towards treatment of carpal tunnel syndrome in adults (defined as patients older than 18 years of age).

These recommendations assume that the patient has reversible mechanical compression of the median nerve based on the diagnostic criteria set forth in the AAOS clinical practice guideline for The Diagnosis of Carpal Tunnel Syndrome. This does not include patients who have nerve damage characterized by irreversible microscopic damage to the nerve ultra-structure. Such cases, understood to exist, without biopsy evidence, have a worse prognosis for recovery with sustained numbness, tingling, paralysis, dyshidrotic changes of the skin, and pain. Diagnostic stratification studies which define preoperative criteria for this division between reversible and irreversible damage were not found. The clinical objective in the more damaged group has lesser expectations and anticipated outcomes by definition.

INCIDENCE AND PREVALENCE

Carpal tunnel syndrome incidence in the United States has been estimated at 1-3 cases per 1000 persons per year. Prevalence is approximately 50 cases per 1000 persons in the general population. 1

BURDEN OF DISEASE

Many Americans experience symptoms of carpal tunnel syndrome and they also expect relief of the condition, which can be accomplished with proper treatment. Untreated or ill-treated carpal tunnel syndrome may worsen and progress to permanent sensory loss and thenar paralysis in some cases.

As carpal tunnel syndrome in the workplace demands attention and as the number of worker's compensation cases are filed increases, the expense for lost productivity and cost of treatment continues to increase. According to the National Institute of Health (NIH), the average lifetime cost of carpal tunnel syndrome, including medical bills and lost time from work, is estimated to be about \$30,000 for each injured worker." Hanrahan et al quote similar estimates by the National Council on Compensation Insurance that estimates the average CTS case costs \$29,000 in Worker's Compensation benefits and medical costs. The Bureau of Labor Statistics reports, as of 2005, the major industry division with highest number of events and exposures is manufacturing. There

were more than 3.8 million visits made to physicians in office-based practices in 2003 because of carpal tunnel syndrome.⁶ According to the Burden of Musculoskeletal Diseases in the United States (2008, p.136), the National Health Interview Survey "is believed to underreport the incidence of injuries" and the Bureau of Labor Statistics only report work related data.⁵

ETIOLOGY

Carpal Tunnel Syndrome (CTS) is among the most common disorders of the upper extremity. It is related to many factors but is thought to be caused by increased pressure on the median nerve in the carpal tunnel at the wrist.⁷

DIAGNOSIS AND TREATMENT

Diagnosis of carpal tunnel syndrome is made on the basis of signs, symptoms, and electro-diagnostic tests, as put forth by the AAOS clinical practice guideline on Diagnosis of Carpal Tunnel Syndrome. ⁸ Appropriate diagnosis is a critical factor to providing treatment.

Treatment for CTS is based on the assumption that final decisions are predicated on patient and physician mutual communication, discussion of available treatment alternatives and procedures applicable to the individual patient. Once the patient has been informed of available alternative non-operative therapies and has discussed these options with his/her physician, an informed decision can be made. Clinician input based on experience with both conservative management and surgical skills increases the probability of identifying patients who will benefit from specific treatment options. Patient compliance with prescribed treatments is also a contributing factor for successful treatment.

RISK FACTORS

Several key co-morbidities and/or human factors potentially increase the risk of developing carpal tunnel syndrome. Primary considerations include advancing age, female gender, and the presence of diabetes and/or obesity. Other risk factors include pregnancy, specific occupations, cumulative and repetitive motion injuries, strong family history, specific medical disorders such as hypothyroidism, autoimmune diseases, rheumatologic diseases, arthritis, renal disease, trauma, anatomic predisposition in the wrist and hand due to shape and size, infectious diseases, and substance abuse. These are all common exclusion criteria in CTS treatment studies and hence these potential risks have not been clearly assessed. ⁹

Persons involved in manual labor in some occupations have a greater incidence and severity of the symptoms.⁷ The relationship between work, co-morbidities and personal factors require good physician judgment, experience with medical evidence and knowledge of the vast occupational literature in assigning and apportioning causation. In many cases, there is no identifiable co-morbidity or causal relationship.

II. METHODS

An AAOS Work Group, consisting of eight physician members, was assembled specifically for the development of this guideline. The Work Group consisted of a diverse group of physician specialists with expertise in treating patients with carpal tunnel syndrome.

PROCESS OVERVIEW

The Work Group, with the assistance of the AAOS staff, began by formulating "simulated recommendations". The simulated recommendations were used to define the scope of the guideline and to refine the literature searches that were conducted. The Work Group, with the assistance of the AAOS medical librarian and staff, completed a systematic review of the relevant literature. Details of the systematic review are provided below.

During the process of developing this guideline, the Work Group participated in a series of conference calls and meetings. When published information of sufficient quality was not available, consensus opinion was employed.

The final draft of the guideline was reviewed by an outside advisory panel (peer review), reviewed internally by the AAOS Board of Directors, Council on Research Quality Assessment and Technology, Board of Councilors, and Board of Specialty Societies (public commentary) and approved by the AAOS Evidence Based Practice Committee, Guideline Oversight and Technology Committee, Council on Research Quality Assessment and Technology, and the Board of Directors.

Peer review of the draft guideline is completed by an outside Peer Review Advisory Panel. Outside Advisory Panels are convened for each AAOS guideline and consist of experts in the guideline's topic area. These experts represent professional societies other than AAOS and are nominated by the guideline Work Group prior to beginning work on the guideline. Non-editorial comments received from each reviewer are documented, reviewed by the Work Group and approved by the Work Group Chairperson. AAOS staff sends each reviewer the approved documentation for his/her comments. For this guideline, thirteen outside peer review organizations were invited to review the draft guideline and all supporting documentation. Seven societies participated in the review of the CTS Treatment guideline draft.

Following response to all reviews, the guideline draft was sent to thirty-one individuals, who were members of the AAOS Board of Directors, Council on Research Quality Assessment and Technology, Board of Councilors, and Board of Specialty Societies for public commentary. Following this period of public commentary, the guideline was submitted for approval.

Within AAOS, multiple iterations of written review were conducted by the participating Work Group, AAOS Guidelines and Technology Oversight Committee, AAOS Evidence Based Practice Committee, and the AAOS Council on Research, Quality Assessment and Technology prior to final approval by the AAOS Board of Directors. The total number of

AAOS reviewers within these governing bodies is fifty-eight. The approval process is documented in Appendix VI.

CONSENSUS DEVELOPMENT

Voting on guideline recommendations will be conducted using a modification of the nominal group technique (NGT), a method previously used in guideline development. Briefly, each member of the guideline Work Group ranked his or her agreement with a guideline recommendation on a scale ranging from 1 to 9 (where 1 is "total disagreement" and 9 is "total agreement"). Consensus is obtained if the number of individuals who do not rate a measure as 7, 8, or 9 is statistically non-significant (as determined using the binomial distribution). Because the number of Work Group members who are allowed to dissent with the recommendation depends on statistical significance, the number of permissible dissenters varies with the size of the work group. The number of permissible dissenters for several work group sizes is given in the table below:

Work Group Size	Number of Permissible Dissenters		
≤3	Not allowed. Statistical significance cannot be obtained		
4-5	0		
6-8	1		
9	1 or 2		

The NGT is conducted by first having members vote on a given recommendation without discussion. If the number of dissenters is "permissible", the recommendation is adopted without further discussion. If the number of dissenters is not permissible, there is further discussion to see whether the disagreement(s) can be resolved. Three rounds of voting are held to attempt to resolve disagreements. If disagreements are not resolved after three voting rounds, no recommendation is adopted.

ARTICLE INCLUSION AND EXCLUSION CRITERIA

Inclusion and exclusion criteria were developed a priori. Articles were retrieved and included only if they met these specific inclusion and exclusion criteria (see Appendix II: Article Inclusions and Exclusions). Supplemental searches were conducted to identify national rates and other information relevant to performance measures.

Work Group members were given the opportunity to supplement the searches of electronic databases with articles not identified by those searches. No additional articles were added by the Work Group for this guideline. Had articles been added, they would have been subjected to the same *a priori* inclusion and exclusion criteria specified in Appendix II.

A total of three hundred thirty-two articles were reviewed for this guideline. Ninety-four articles met all *a priori* inclusion criteria. Two hundred and thirty eight articles were excluded for various reasons. Tracking these numbers in the flowcharts is not possible because a study could be included in more than one flowchart (i.e. some of the surgical studies were included in the infection flowchart). These numbers can be verified using the evidence tables and counting the references in the technical report. The flowcharts in Appendix II: Article Inclusions and Exclusions illustrate the number of articles retrieved for specific recommendations as well as the number of articles used to update systematic reviews.

For all recommendations except recommendation 3, we included only studies that diagnosed patients with a combination of electro-diagnostic tests and signs and symptoms. For recommendation 3, which addresses workplace issues, we required only that patients be diagnosed with signs and symptoms (see Appendix II: Article Inclusions and Exclusions). We relaxed the inclusion criteria for studies addressing CTS in the workplace because these patients are typically symptomatic and rarely receive electro-diagnostic tests to confirm their diagnosis. Even though we relaxed our inclusion criteria we were unable to find relevant literature which, to us, indicates a critical need for future research in this area.

We did not search for, or include, all available evidence. Wherever appropriate, we searched for and included the best available evidence. Hence, if Level II evidence was available, we did not search for or include Level III evidence or lower unless there was very little Level II evidence, and a great deal of Level III evidence.

Our analyses focused on patient-oriented outcome measures. These measures are defined in clinical research as "outcomes that matter to patients including reduced morbidity, reduced mortality, symptom improvement, or improving patients' quality of life". ¹¹ By critically focusing on patient-oriented outcomes, the recommendations in this guideline are expected to improve overall patient care in the treatment of carpal tunnel syndrome.

LITERATURE SEARCHES

We searched four electronic databases, MEDLINE, EMBASE, CINAHL and the Cochrane database of systematic reviews, to identify literature for this guideline. Search strategies were reviewed by the work group prior to conducting the searches. A list of the electronic databases we searched and the search strategies we used are provided in Appendix I: Literature Searches. All literature searches were supplemented with manual screening of bibliographies in publications accepted for inclusion into the evidence base. In addition, the bibliographies of recent review articles were searched for potentially relevant citations. All included articles met the specified a priori inclusion/exclusion criteria.

ASSIGNING A LEVEL OF EVIDENCE

The quality of evidence was rated using the evidence hierarchy shown in Appendix III: Rating Evidence Quality. A complete description of the hierarchy is included in the AAOS Evidence Report for this guideline. This hierarchy is also on the American Academy of Orthopaedic Surgeons (AAOS) website at: http://www.aaos.org/Research/Committee/Evidence/loetable1.pdf

DATA EXTRACTION

Six reviewers independently completed data extraction for all studies. Evidence tables were constructed to summarize the best evidence pertaining to each recommendation and all evidence can be found in the accompanying Evidence Report ⁹ to this guideline.

GRADING THE RECOMMENDATIONS

Each guideline recommendation was graded using the following system:

- A: Good evidence (Level I Studies with consistent findings) for or against recommending intervention.
- B: Fair evidence (Level II or III Studies with consistent findings) for or against recommending intervention.
- C: Poor quality evidence (Level IV or V) for or against recommending intervention.
- I: There is insufficient or conflicting evidence not allowing a recommendation for or against intervention.

The Committee used the following language in constructing the recommendations:

We recommend Treatment X: (for Grade A recommendations)
We suggest Treatment X: (for Grade B recommendations)
Treatment X is an option: (for Grade C recommendations)

These definitions^{12, 13} help clarify the intent of the Work Group by reflecting the assessment of the importance of adherence to the recommendation based on the grade of the recommendation

STATISTICAL METHODS

The statistical analyses performed help compare the treatment options available to patients with carpal tunnel syndrome. In order to assess specific treatments, comparisons were made between similar populations of patients receiving the treatment to patients receiving a control, placebo, or a second treatment. The goal of most treatment comparisons is to demonstrate that a treatment has a significant effect or that there is a significant difference between two treatments.

Small sample sizes in clinical trials present serious concerns because a lack of statistical power means that small but clinically important differences may go undetected. We calculated the minimal detectable difference to determine if a study was sufficiently powered for the given outcome. In our power calculations, we used 80% power, 95% confidence intervals and the number of patients per group. This allowed calculation of the minimal detectable effect size which was compared to the calculated effect size to

determine if the study had enough power to detect the observed effect. If the trial was found to lack sufficient power for a given outcome, its results were taken as inconclusive. Power calculations were performed using G Power 3 (Version 3.0.5). Results are listed in the Evidence Report 9 and Evidence Tables. 15

For recommendations one through eight in this guideline, several measures of association including the odds ratio (OR) and the natural log of the odds ratio (log OR) were used to compare treatments. In addition, the standardized mean difference (SMD) was used for computing standardized measures of effect size. Effect sizes were calculated when applicable; OR and log OR for dichotomous data and the SMD for continuous data. The larger the OR is, the larger the effect size. The SMD can be evaluated as follows: 0.2 for a small effect, 0.5 for a moderate effect and 0.8 for a large effect. ¹⁶

Studies had to have treatments, outcome measures, and durations of follow up in common to perform meta-analysis of the data. Given the paucity and heterogeneity of the data for specific recommendations, we did not apply formal meta-analytic techniques in all circumstances. Log OR and Cohen's h were computed for dichotomous outcome measures and SMD was computed for continuous outcome measures that were pooled for meta-analyses. When the event rate was zero for dichotomous outcome measures, a continuity correction was added. The Log OR and SMD values were then meta-analyzed using standard DerSimonian and Laird random effects model meta-analysis. When possible, effect sizes were pooled across different studies, and heterogeneity was assessed with the I-squared statistic. Summary statistics were presented when heterogeneity was less than 50%. All meta-analyses and effect size calculations were performed using STATA 10.0 (StataCorp LP, College Station, Texas) and the "metan" command.

Recommendation nine addressed the applicability of various instruments for the evaluation of carpal tunnel syndrome treatment in patients. Instruments are generally evaluated using eight key component areas: appropriateness, reliability, validity, responsiveness, precision, interpretability, acceptability, and feasibility. For this evaluation, we did not assess appropriateness, precision, interpretability, acceptability or feasibility. The physician should consider whether the instrument used was appropriate to measure CTS outcomes, that it contained the appropriate number of distinctions with regard to the dimensions being measured for precision and that the instruments were generally acceptable and feasible for use in the identified patient population. An overall summary of the properties assessed in each instrument (by primary study) is illustrated in the Evidence Tables. ¹⁵

Reliability, validity, and responsiveness were the three primary key components addressed in the studies for this recommendation. Reliable instruments are internally consistent and internal consistency is commonly measured by Cronbach's alpha. This statistic measures how comparable the results of the instrument would be if the instrument were split into two versions or the average level of agreement of all the possible ways of performing split-half tests.¹⁸

Validity was quantified using Spearman and Pearson correlation coefficients. Instruments with similar concepts should have large correlations and instruments with different

concepts should have small correlations. A correlation coefficient of 0.5 and above is a large correlation, indicating two converging instruments, and a correlation coefficient below 0.5 is a smaller correlation, indicating two diverging instruments. ¹⁹ A negative correlation indicates two instruments that score in the opposite direction.

Responsiveness was measured using the standardized response mean (SRM). The SRM is expressed as the change score divided by the standard deviation of the change score. A standardized response mean of 0.2 is indicative of a small change, 0.5 a medium change, and 0.8 a large change. ^{19, 20}

REVISION PLANS

This guideline represents a cross-sectional view of current intervention methods and will become outdated when more sophisticated tests, more objective assessments and more rigorous differential diagnosis are possible. Linkage to other disorders, genetic diagnosis, and occupational and human factors literature will contribute to our understanding of the early stages of this condition and the means of differential treatment.

Because of the high profile of CTS in the workplace and the high level of interest in this topic, the guideline will be revised in accordance with changing practice, rapidly emerging opinion, new technology, and new evidence. It is anticipated that this guideline will be revised in 2011.

III. RECOMMENDATIONS

RECOMMENDATION 1

A course of non-operative treatment is an option in patients diagnosed with carpal tunnel syndrome. Early surgery is an option when there is clinical evidence of median nerve denervation or the patient elects to proceed directly to surgical treatment.

(Grade C, Level V)

Rationale

Data were extracted from three systematic reviews and twenty-three randomized controlled or controlled trials for evidence to support this recommendation. The literature found supported the effectiveness of non-operative treatment over placebo. Data were not found that clearly identified when non-operative treatment should be considered the only option, nor were studies found in which non-operative treatment was clearly shown to be completely ineffective and therefore contraindicated.

Studies of carpal tunnel syndrome often included denervation as an indication for surgery, and a relative contraindication for non-surgical treatment, so such cases were not studied systematically. Consequently, it was not possible to make a Grade A or B recommendation. Therefore, this guideline recommendation is, of necessity, based upon expert opinion.

See Evidence Tables 1-21 and Evidence Report page 12.9,15

We suggest another non-operative treatment or surgery when the current treatment fails to resolve the symptoms within 2 weeks to 7 weeks.

(Grade B, Level I and II)

Rationale

Considerable evidence exists that suggests patients benefit from a variety of non-operative treatment and surgical options for carpal tunnel syndrome. Although the data did not report the minimum time for effectiveness, an analysis of the level I and II data reviewed for Recommendations 4a-c suggested that all effective or potentially effective non-operative treatments (local steroid injections, splinting, oral steroids and ultrasound) for carpal tunnel syndrome have a measurable effect on symptoms within two to seven weeks of the initiation of treatment. If a treatment is not effective in reducing symptoms within that time frame, then consideration should be given to trying a different one, assuming, of course, that the diagnosis of carpal tunnel syndrome is not in doubt.

Because this recommendation considers a variety of non-operative treatments, the levels of evidence varied. More level II evidence exists than level I evidence; hence the grade of recommendation is based on consistent level II evidence.

See Evidence Tables 1-21 and Evidence Report page 13.9,15

We do not have sufficient evidence to provide specific treatment recommendations for carpal tunnel syndrome when found in association with the following conditions: diabetes mellitus, coexistent cervical radiculopathy, hypothyroidism, polyneuropathy, pregnancy, rheumatoid arthritis, and carpal tunnel syndrome in the workplace.

(Inconclusive, No evidence found)

Rationale

Despite an exhaustive review of the literature, there was insufficient evidence to make conclusions about these conditions and carpal tunnel syndrome in the workplace. These potentially treatable medical conditions are common exclusion criteria from controlled trials. This makes it difficult to make specific recommendations for how to treat such patients.

See Evidence Tables 1-21 and Evidence Report pages 12-15. 9,15

Local steroid injection or splinting is suggested when treating patients with carpal tunnel syndrome, before considering surgery.

(Grade B, Level I and II)

Rationale

Local steroid injection and splinting are effective in treating carpal tunnel syndrome. Splinting was effective at 2, 4, and 12 weeks in reducing symptoms and improving functional status. ^{84, 93} No conclusion could be drawn at the 6 month time point because the studies were underpowered.

Steroid injections are also effective for treating carpal tunnel syndrome. Patient satisfaction (2 weeks ²³), clinical improvement (4 weeks ^{44, 77} 8 weeks, ¹¹⁶ 12 weeks ¹¹⁶), symptoms (2 weeks, ⁵⁸ 4 months, ²⁷ 6 months ¹⁰⁸), function (3 months ¹⁰⁸), and pain (8 weeks ⁵⁸) were shown to improve after cortisone injections.

Patients with more severe or prolonged CTS, however defined, may not benefit from prolonged, non-operative treatment. Trials of non-operative treatment are suggested for the treating physician and should show remission as described in the recommendations above at the intervals indicated.

See Evidence Tables 1- 21 and Evidence Report pages 16-61, figures 1-6, & 37-41. 9,15

RECOMMENDATION 4B

Oral steroids or ultrasound are options when treating patients with carpal tunnel syndrome.

(Grade C, Level II).

Rationale

Oral steroid treatment was effective in the treatment of carpal tunnel syndrome.^{63, 84} However, the evidence suggested that local steroid injection is more effective than oral steroids.⁷⁷ Since the evidence supports other more effective treatments, the Work Group downgraded the recommendation about oral steroids to Grade C, "optional".

Ultrasound was also shown to be effective in the treatment of carpal tunnel syndrome in two studies. ^{28,84} One of the studies ²⁸ however, compared ultrasound to laser treatment, an unproven modality, rather than to a control. Hence, there was only one level II study supporting ultrasound. Based on this methodological flaw, the Work Group chose to downgrade this recommendation on ultrasound to Grade C, "optional".

See Evidence Tables 1-21 and Evidence Report pages 16-61, figures 12-15, 23, & 24.9,15

We recommend carpal tunnel release as treatment for carpal tunnel syndrome.

(Grade A, Level I)

Rationale

Level I evidence demonstrates that surgical release of the flexor retinaculum is an extremely effective treatment for patients with carpal tunnel syndrome. The evaluation of operative versus non-operative treatment of carpal tunnel syndrome demonstrated the effectiveness of the surgical treatment.

These recommendations assume that the patient has reversible mechanical compression of the median nerve based on the diagnostic criteria set forth in the AAOS clinical practice guideline for The Diagnosis of Carpal Tunnel Syndrome. This does not include patients who have nerve damage characterized by irreversible microscopic damage to the nerve ultra-structure. Such cases, understood to exist, without biopsy evidence, have a worse prognosis for recovery with sustained numbness, tingling, paralysis, dyshidrotic changes of the skin, and pain. Diagnostic stratification studies which define preoperative criteria for this division between reversible and irreversible damage were not found. The clinical objective in the more damaged group has lesser expectations and anticipated outcomes by definition.

See Evidence Tables 1-21 and Evidence Report pages 62-66, figures 53-58. 9,15

Heat therapy is not among the options that should be used to treat patients with carpal tunnel syndrome.

(Grade C, Level II)

Rationale

Heat therapy was less effective than placebo control in treating carpal tunnel syndrome.⁸¹ The grade of recommendation is based on a single study therefore, it was downgraded to Grade C, "optional".

See Evidence Tables 1-21 and Evidence Report page 43, figure 30.9,15

RECOMMENDATION 4E

The following treatments carry no recommendation for or against their use: activity modifications, acupuncture, cognitive behavioral therapy, cold laser, diuretics, exercise, electric stimulation, fitness, graston instrument, iontophoresis, laser, stretching, massage therapy, magnet therapy, manipulation, medications (including anticonvulsants, antidepressants and NSAIDs), nutritional supplements, phonophoresis, smoking cessation, systemic steroid injection, therapeutic touch, vitamin B6 (pyridoxine), weight reduction, yoga.

(Inconclusive, Level II and V).

Rationale

Despite an extensive review of the literature, there was insufficient evidence to make conclusions about these modalities. For some treatments, there were simply no studies that met the inclusion criteria. For others, the studies had too little statistical power to allow for meaningful conclusions. Still other studies were downgraded from a higher grade of recommendation because their applicability was questioned. Consequently, we are unable to make recommendations for or against the use of these treatments.

One study compared the Graston Instrument to manual therapy.³⁷ The applicability of this study was questioned because the Graston instrument was compared to an unproven alternative treatment. This was the only study looking at the Graston instrument that met the inclusion criteria. The grade of recommendation was downgraded because the evidence was inconclusive.

One systematic review⁸⁴ examined the comparison of Vitamin B (pyridoxine) to placebo. The applicability of the outcome measure was questioned because it was not considered to be critical to determining whether Vitamin B was beneficial in the treatment of CTS. The grade of recommendation was downgraded because the evidence was inconclusive.

All of these modalities require further investigation in appropriately designed studies to determine their efficacy in the treatment of carpal tunnel syndrome.

See Evidence Tables 1- 21 and Evidence Report pages 16-61. 9,15

Table A. Recommendation 4e Summary of Treatment Evidence

Reasoning for Insufficient Evidence

We recommend surgical treatment of carpal tunnel syndrome by complete division of the flexor retinaculum regardless of the specific surgical technique.

(Grade A, Level I and II).

Rationale

Complete division of the flexor retinaculum is an effective method for treating CTS. Two systematic reviews ^{97,107} and six randomized controlled trials ^{26,32,39,95,96,115} examined comparisons between open carpal tunnel release, endoscopic carpal tunnel release, or minimal incision carpal tunnel release. Several patient-oriented outcome measures, including symptom severity and functional status at 52 weeks post-operatively, residual pain at 12 weeks post-operatively, reversible nerve damage, return to work and wound-related complications, were evaluated using meta-analytic techniques to compare open release and endoscopic release. Endoscopic release was favored in residual pain at 12 weeks post-operatively, return to work time, and wound related complications. Open release was favored when reversible nerve damage was the outcome compared. No difference in the techniques was found in symptom severity or functional status at 52 weeks, complications, and infections.

In addition, minimal incision release was compared to open or endoscopic release in Level I studies. When compared to open release, minimal incision was favored in symptom severity, functional status, and scar tenderness. When compared to endoscopic release, minimal incision was favored when pain at two or four weeks was the outcome measure.

The Work Group discussed the studies and agreed that not all relevant outcomes were available, addressed, and/or analyzed by the evidence comparing the various surgical techniques. Nevertheless, Level I and Level II evidence clearly indicates the effectiveness of complete division of the flexor retinaculum, regardless of surgical technique, as a treatment for CTS.

See Evidence Tables 23-37 and Evidence Report pages 68-87, figures 59-81. 9,15

We suggest that surgeons not routinely use the following procedures when performing carpal tunnel release:

skin nerve preservation (Grade B, Level I)

epineurotomy (Grade C, Level II)

The following procedures carry no recommendation for or against their use: flexor retinaculum lengthening, internal neurolysis, tenosynovectomy, ulnar bursa preservation.

(Inconclusive, Level II and V)

Rationale

A single Level I study ¹⁰¹ evaluated the effect of preserving cutaneous nerves in the path of a skin incision made in the customary location for a carpal tunnel release. Preservation was compared to a standard approach to making a skin incision, which did not seek to preserve any nerve branches encountered as the wound was deepened down to the palmar fascia. The Patient Evaluation Measure (PEM) indicated a slight advantage in favor of the standard approach at the three-month assessment. The PEM is a broader evaluation of outcome than the VAS suggesting that the advantages for a standard carpal tunnel release incision refer to a domain other than pain.

Epineurotomy was studied in a systematic review and in a single Level II study. In the systematic review ⁹⁷ the outcome was described as "overall improvement" at 12 months and, in the single Level II study, ³⁴ the outcomes were "nocturnal pain" and "paraesthesia" at three months following surgery. Both studies indicated a mild effect favoring no epineurotomy.

Tenosynovectomy and internal neurolysis were compared in a systematic review ⁹⁷ and the data were inconclusive. Lengthening of the flexor retinaculum was studied in a Level I study ⁴⁹ that used the Boston Carpal Tunnel Questionnaire as the outcome measure. The results were inconclusive because the study had too little power to allow for statistically meaningful comparison. A single Level I study ⁵³ examining ulnar bursa preservation, with VAS and PEM as the outcome measures at 8 weeks, also had too little power to allow for meaningful statistical comparisons. The study was therefore inconclusive.

See Evidence Tables 23-37 and Evidence Report pages 88-95, figures 82-88. 9,15

The physician has the option of prescribing pre-operative antibiotics for carpal tunnel surgery.

(Grade C, Level III)

Rationale

Our searches indicated that the current literature rarely reports whether pre-operative antibiotic treatment was used in carpal tunnel release. Of forty-five studies analyzed for this recommendation, forty-four did not report whether pre-operative antibiotics were used. The study that did report antibiotic use reported that 6.03% of patients developed a post-operative infection, even though all patients received antibiotics.

An examination of the various trials addressing carpal tunnel syndrome treatment did not provide insight on whether there are conditions or comorbidities that predispose patients to post-surgical infection. Patients with diabetes mellitus, for example, were excluded from the trials. A single Level IV study looked at rates of post-operative infections in persons with and without diabetes and found that the rate was similar in the two groups.

See Evidence Tables 38-41 and Evidence Report pages 96-100. 9,15

We suggest that the wrist not be immobilized postoperatively after routine carpal tunnel surgery.

(Grade B, Level II)

We make no recommendation for or against the use of postoperative rehabilitation. (Inconclusive, Level II).

Rationale

The wrist should not be immobilized postoperatively after routine carpal tunnel release. Post-operative splinting for longer than two weeks did not offer any specific benefit in terms of grip or lateral pinch strength, bowstringing, complication rates, subjective outcome and patient satisfaction. 38,43,52,78

Clinicians may wish to provide protection for the wrist in a working environment or for temporary protection. However, the evidence does not provide objective criteria for these situations. Clinicians should be aware of the detrimental affects including adhesion formation, stiffness and prevention of nerve and tendon movement which may compromise the carpal tunnel release results in achieving another objective such as early release to work.

For postoperative rehabilitation, one study examined supervised hand therapy ⁹⁴. The applicability of the outcome measure (return to work) was questioned because it was not considered to be critical to determining whether supervised hand therapy was beneficial to postoperative rehabilitation. The grade of recommendation was downgraded because the evidence was inconclusive.

There were no included studies that looked at work hardening and the role of various modalities for post-operative carpal tunnel management. The role of supervised therapy after carpal tunnel release in the work-related population will need further evaluation to determine if there is any advantage to work hardening, work simulation, or routine strengthening.

See Evidence Tables 42-51 and Evidence Report pages 101-110, figures 92-101. 9,15

We suggest physicians use one or more of the following instruments when assessing patients' responses to CTS treatment for research:

Boston Carpal Tunnel Questionnaire (disease-specific)

DASH – Disabilities of the arm, shoulder, and hand (region-specific; upper limb)

MHQ – Michigan Hand Outcomes Questionnaire (region-specific; hand/wrist)

PEM (region-specific; hand)

SF-12 or SF-36 Short Form Health Survey (generic; physical health component for global health impact)

(Grade B, Level I, II, and III)

Rationale

All measurement instruments, whether they are aimed at diagnosis, evaluation of disease activity or outcome, must be judged on their key psychometric characteristics: reliability, validity, interpretability and responsiveness. Reliability was generally measured in these studies by assessing the internal consistency and reproducibility of the study.

Per Jenkinson ⁶⁷, "validity is assessed in relation to a specific purpose and setting." Validity is established statistically for an instrument by measuring construct validity, convergent and divergent validity, and/or criterion validity. Instruments having construct validity are summarized in the table below.

Convergent and divergent validity measures can be found in the Evidence Tables (Tables 69-85). More information concerning interpretation of these measures can be found in the Evidence Report but an inclusive discussion is beyond the scope of the guideline. These values were not graphed and were provided to illustrate the direction and magnitude of relationships. Criterion validity was not summarized.

While adequate reliability and validity are concepts that are, for the most part, clear to clinicians, the capacity for interpretability and responsiveness may be less familiar. Interpretability refers to the fundamental meaning of the measure. Instruments that encompass items that are meaningful to patients and/or clinicians will have good interpretability and users can easily understand the meaning of these measures. Few studies measured interpretability therefore they are not summarized in the table below.

Responsive measures reflect small changes in a given condition. This may be important where subtle differences could be clinically important. Responsive measures are helpful in the planning of trials where the objective may be to demonstrate a small difference between, for example, treatments.

Generally speaking, generic measures, like the Short Form 36 (SF-36), look at a broadly based assessment of health and, as a result may not be very responsive to changes in status related to a relatively minor condition such as CTS.

Disease-specific instruments such as the Boston Carpal Tunnel Questionnaire (BCTQ) are most responsive. ⁹ The BCTQ shows excellent responsiveness for the measurement of disease activity in CTS. Wherever possible the full instrument should be used because this gives the most comprehensive evaluation of both function and symptoms in CTS without any loss of responsiveness. The subscales of this instrument also have satisfactory responsiveness ⁹ but give a more narrow view of disease activity. The BCTQ is fully validated in the treatment of carpal tunnel syndrome.

The region-specific instrument, The Disabilities of the Arm, Shoulder and Hand (DASH) was moderate to highly responsive ⁹ and the Michigan Hand Outcomes Questionnaire (MHQ) was highly responsive in three of five subscales .⁹

The Patient Evaluation Measure (PEM), (MHQ) and DASH are more broadly based region-specific instruments that can be considered to be responsive for the evaluation of CTS. The responsiveness of the DASH is slightly below the acceptable threshold (standardized response mean (SRM) = 0.80) but should be considered if the goal of the evaluation is a focus on disability because it has been evaluated in three key domains: internal consistency, reproducibility and responsiveness.

See Evidence Tables 52-101 and Evidence Report pages 111-125, figures 102-120. 9,15

Table B. Psychometric Properties of Instruments

Instrument	Internal Consistency (Reliability)	Reproducibility (Reliability)	Construct Validity (Validity)	Responsiveness (SRM)
BCTQ-Total				>0.8
BCTQ-SSS	X	X	X	>0.8
BCTQ-FSS	X	X	X	>0.8
AIMS2 subscales*				0.06 - 1.72
DASH	X	X	X	0.76
MHQ subscales*				0.5 – 1.1
PEM		X	X	>0.8
VAS			X	0.51
SF-36 subscales*				0 - 0.86
SF-12 subscales*				0.08 - 0.58

^{*} See the Evidence Report for CTS Treatment for responsiveness of individual subscales.9

FUTURE RESEARCH

Although we make every effort to find studies of the highest quality, such evidence is not readily available for carpal tunnel syndrome treatment at this time. This guideline has been hindered by a relative lack of power in the studies even though these studies were of Level I and II evidence. The recommendations of this guideline therefore depend to some degree on lesser evidence, including expert opinion.

To achieve a high-quality literature base, academic authors and scientists should invest their time and effort in studies designed to avoid bias (e.g., blinded and properly randomized controlled trials of sufficient power to address the outcome of interest). Future studies should, from the onset, be based on improved study design that includes *a priori* power calculations. Risk stratification studies are also needed to detect when antibiotics might be justified on the basis of co-morbidities and co-interventions.

We recognize that the issue of carpal tunnel syndrome in the workplace is important. Studies identified by the literature search commonly analyze risk, prevalence, and predictability of carpal tunnel syndrome in specific job categories but good evidence to address the effectiveness of workplace modifications was not available. Working patients, payors, and physicians clearly lack the evidence base to determine "best options". Physicians and patients must first decide the desired outcome. Should the goal be permanent modification of activities for the worker or proceed to surgery and return to normal activities? Future research must rigorously address this subpopulation to determine if activity modification will result in positive outcomes such as ultimately avoiding surgery.

REFERENCES

- (1) Ashworth NL, Carpal Tunnel Syndrome, Retrieved 03/14/2007, http://www.emedicine.com/pmr/topic21.htm, Last updated November 30, 2006.
- (2) National Institute of Neurological Diseases and Stroke (NINDS), http://www.ninds.nih.gov/disorders/carpal_tunnel/detail_carpal_tunnel.htm, Last updated February 12, 2007.
- (3) Hanrahan LP., Higgins D., Anderson H., Smith M. Wisconsin occupational carpal tunnel syndrome surveillance: the incidence of surgically treated cases. Wisconsin Medical Journal 1993; 92 (12): 695-689.
- (4) US Department of Labor Bureau of Labor Statistics, Retrieved on 10/18/2007, http://www.bls.gov/news.release/osh2.t05.htm, Last updated 11/2006.
- (5) American Academy of Orthopeadic Surgeons (2008). The Burden of Musculoskeletal Diseases in the United States, AAOS
- (6) National Center for Health Statistics, National Ambulatory Medical Care Survey 2000, data was extracted using all three possible reason-for-visit codes identified; data extracted and compiled by the AAOS Department of Research and Scientific Affairs in 2003. http://orthoinfo.aaos.org/topic.cfm?topic=A00130#A00130_R2_anchor
- (7) Fuller DA, Carpal Tunnel Syndrome, Retrieved 03/14/2007 http://www.emedicine.com/orthoped/topic455.htm, Last updated July 2, 2004
- (8) American Academy of Orthopaedic Surgeons clinical practice guideline on Diagnosis of Carpal Tunnel Syndrome (2007), http://www.aaos.org/Research/guidelines/CTS_guideline.pdf
- (9) Treatment of Carpal Tunnel Syndrome Evidence Report, http://www.aaos.org/research/guidelines/CTSTreatmentEvidenceReport2.pdf
- (10) Murphy MK, Black LA, Lamping, DL, McKee CM, Sanderson, CFB, Askham J, et al. Consensus development methods, and their use in clinical guideline development Health Technol. Assessment 1998; 2(3).
- (11) Ebell MH, Siwek J, Weiss BD, Woolf SH, Susman J, Ewigman B, et al. Strength of recommendation taxonomy (SORT): a patient-centered approach to grading evidence in the medical literature. J Am Board Fam Pract 2004 Jan;17(1):59-67.
- (12) Guyatt G, Gutterman D, Baumann MH, Addrizzo-Harris D, Hylek EM, Phillips B, et al. Grading strength of recommendations and quality of evidence in clinical practice guidelines: report from an American College of Chest Physicians task force. Chest 2006 Jan;129(1):174-81.

- (13) Classifying recommendations for clinical practice guidelines. Pediatrics 2004 Sep;114(3):874-7.
- (14) Faul F, Erdfelder E, Lang AG, Buchner A. G*Power 3: a flexible statistical power analysis program for the social, behavioral, and biomedical sciences. Behav Res Methods 2007 May;39(2):175-91.
- (15) Treatment of Carpal Tunnel Syndrome Evidence Tables, http://www.aaos.org/research/guidelines/CTSTreatmentEvidenceTables.pdf
- (16) Higgins JP, Thompson SG. Quantifying heterogeneity in a meta-analysis. Stat Med 2002 Jun 15;21(11):1539-58.
- (17) DerSimonian R, Laird N. Meta-analysis in clinical trials. Controlled Clinical Trials, 1986; 7: 177-188.
- (18) Fitzpatrick R, Davey C, Buxton MJ, Jones DR. Evaluating patient-based outcome measures for use in clinical trials. Health Technol Assess 1998;2(14):i-74.
- (19) Cohen J. Statistical power analysis for the behavioural sciences. New York: Academic Press; 1977.
- (20) Kazis LE, Anderson JJ, Meenan RF. Effect sizes for interpreting changes in health status. Med Care 1989 Mar;27(3 Suppl):S178-S189.

REFERENCES TO INCLUDED ARTICLES

- (21) Amadio PC, Silverstein MD, Ilstrup DM, Schleck CD, Jensen LM. Outcome after Colles fracture: the relative responsiveness of three questionnaires and physical examination measures. J Hand Surg [Am] 1996 Sep;21(5):781-7.
- (22) Amadio PC, Silverstein MD, Ilstrup DM, Schleck CD, Jensen LM. Outcome assessment for carpal tunnel surgery: the relative responsiveness of generic, arthritis-specific, disease-specific, and physical examination measures. J Hand Surg [Am] 1996 May;21(3):338-46.
- (23) Armstrong T, Devor W, Borschel L, Contreras R. Intracarpal steroid injection is safe and effective for short-term management of carpal tunnel syndrome. Muscle Nerve 2004 Jan;29(1):82-8.
- (24) Atroshi I, Breidenbach WC, McCabe SJ. Assessment of the carpal tunnel outcome instrument in patients with nerve-compression symptoms. J Hand Surg [Am] 1997 Mar;22(2):222-7.
- (25) Atroshi I, Johnsson R, Ornstein E. Endoscopic carpal tunnel release: prospective assessment of 255 consecutive cases. J Hand Surg [Br] 1997 Feb;22(1):42-7.

- (26) 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 Jun 24;332(7556):1473.
- (27) Aygul R, Ulvi H, Karatay S, Deniz O, Varoglu AO. Determination of sensitive electrophysiologic parameters at follow-up of different steroid treatments of carpal tunnel syndrome. J Clin Neurophysiol 2005 Jun;22(3):222-30.
- (28) Bakhtiary AH, Rashidy-Pour A. Ultrasound and laser therapy in the treatment of carpal tunnel syndrome. Aust J Physiother 2004;50(3):147-51.
- (29) Baysal O, Altay Z, Ozcan C, Ertem K, Yologlu S, Kayhan A. Comparison of three conservative treatment protocols in carpal tunnel syndrome. Int J Clin Pract 2006 Jul;60(7):820-8.
- (30) Beaton DE, Katz JN, Fossel AH, Wright JG, Tarasuk V, Bombardier C. Measuring the whole or the parts? Validity, reliability, and responsiveness of the Disabilities of the Arm, Shoulder and Hand outcome measure in different regions of the upper extremity. J Hand Ther 2001 Apr;14(2):128-46.
- (31) Bessette L, Sangha O, Kuntz KM, Keller RB, Lew RA, Fossel AH, et al. Comparative responsiveness of generic versus disease-specific and weighted versus unweighted health status measures in carpal tunnel syndrome. Med Care 1998 Apr;36(4):491-502.
- (32) Bhattacharya R, Birdsall PD, Finn P, Stothard J. A randomized controlled trial of knifelight and open carpal tunnel release. J Hand Surg [Br] 2004 Apr;29(2):113-5.
- (33) Blankfield RP, Sulzmann C, Fradley LG, Tapolyai AA, Zyzanski SJ. Therapeutic touch in the treatment of carpal tunnel syndrome. J Am Board Fam Pract 2001 Sep;14(5):335-42.
- (34) Borisch N, Haussmann P. Neurophysiological recovery after open carpal tunnel decompression: comparison of simple decompression and decompression with epineurotomy. J Hand Surg [Br] 2003 Oct;28(5):450-4.
- (35) Brazier JE, Harper R, Jones NM, O'Cathain A, Thomas KJ, Usherwood T, et al. Validating the SF-36 health survey questionnaire: new outcome measure for primary care. BMJ 1992 Jul 18;305(6846):160-4.
- (36) Brief R, Brief LP. Endoscopic carpal tunnel release: report of 146 cases. Mt Sinai J Med 2000 Sep;67(4):274-7.
- (37) Burke J, Buchberger DJ, Carey-Loghmani MT, Dougherty PE, Greco DS, Dishman JD. A pilot study comparing two manual therapy interventions for carpal tunnel syndrome. J Manipulative Physiol Ther 2007 Jan;30(1):50-61.

- (38) Bury TF, Akelman E, Weiss AP. Prospective, randomized trial of splinting after carpal tunnel release. Ann Plast Surg 1995 Jul;35(1):19-22.
- (39) Cellocco P, Rossi C, Bizzarri F, Patrizio L, Costanzo G. Mini-open blind procedure versus limited open technique for carpal tunnel release: a 30-month follow-up study. J Hand Surg [Am] 2005 May;30(3):493-9.
- (40) Chow JC. The Chow technique of endoscopic release of the carpal ligament for carpal tunnel syndrome: four years of clinical results. Arthroscopy 1993;9(3):301-14.
- (41) Chung KC, Pillsbury MS, Walters MR, Hayward RA. Reliability and validity testing of the Michigan Hand Outcomes Questionnaire. J Hand Surg [Am] 1998 Jul;23(4):575-87.
- (42) Cobb TK, Amadio PC, Leatherwood DF, Schleck CD, Ilstrup DM. Outcome of reoperation for carpal tunnel syndrome. J Hand Surg [Am] 1996 May;21(3):347-56.
- (43) Cook AC, Szabo RM, Birkholz SW, King EF. Early mobilization following carpal tunnel release. A prospective randomized study. J Hand Surg [Br] 1995 Apr;20(2):228-30.
- (44) Dammers JW, Roos Y, Veering MM, Vermeulen M. Injection with methylprednisolone in patients with the carpal tunnel syndrome: a randomised double blind trial testing three different doses. J Neurol 2006 May;253(5):574-7.
- (45) Davies BW, Pennington GA, Fritz AM. Two-portal endoscopic carpal tunnel release: an outcome analysis of 333 hands. Ann Plast Surg 1998 May;40(5):542-8.
- (46) De SL, De KR, Degreef I, Debeer P. Responsiveness of the Dutch version of the DASH as an outcome measure for carpal tunnel syndrome. J Hand Surg [Br] 2007 Feb;32(1):74-6.
- (47) Demirci S, Kutluhan S, Koyuncuoglu HR, Kerman M, Heybeli N, Akkus S, et al. Comparison of open carpal tunnel release and local steroid treatment outcomes in idiopathic carpal tunnel syndrome. Rheumatol Int 2002 May;22(1):33-7.
- (48) Dias JJ, Bhowal B, Wildin CJ, Thompson JR. Assessing the outcome of disorders of the hand. Is the patient evaluation measure reliable, valid, responsive and without bias? J Bone Joint Surg Br 2001 Mar;83(2):235-40.
- (49) Dias JJ, Bhowal B, Wildin CJ, Thompson JR. Carpal tunnel decompression. Is lengthening of the flexor retinaculum better than simple division? J Hand Surg [Br] 2004 Jun;29(3):271-6.

- (50) Evcik D, Kavuncu V, Cakir T, Subasi V, Yaman M. Laser therapy in the treatment of carpal tunnel syndrome: a randomized controlled trial. Photomed Laser Surg 2007 Feb;25(1):34-9.
- (51) Fagan DJ, Evans A, Ghandour A, Prabhkaran P, Clay NR. A controlled clinical trial of postoperative hand elevation at home following day-case surgery. J Hand Surg [Br] 2004 Oct;29(5):458-60.
- (52) Finsen V, Andersen K, Russwurm H. No advantage from splinting the wrist after open carpal tunnel release. A randomized study of 82 wrists. Acta Orthop Scand 1999 Jun;70(3):288-92.
- (53) Forward DP, Singh AK, Lawrence TM, Sithole JS, Davis TR, Oni JA. Preservation of the ulnar bursa within the carpal tunnel: does it improve the outcome of carpal tunnel surgery? A randomized, controlled trial. J Bone Joint Surg Am 2006 Nov;88(11):2432-8.
- (54) Forward DP, Sithole JS, Davis TR. The internal consistency and validity of the patient evaluation measure for outcomes assessment in distal radius fractures. J Hand Surg [Br] 2007 Jun;32(3):262-7.
- (55) Gainer JV, Jr., Nugent GR. Carpal tunnel syndrome: report of 430 operations. South Med J 1977 Mar;70(3):325-8.
- (56) Gay RE, Amadio PC, Johnson JC. Comparative responsiveness of the disabilities of the arm, shoulder, and hand, the carpal tunnel questionnaire, and the SF-36 to clinical change after carpal tunnel release. J Hand Surg [Am] 2003 Mar;28(2):250-4.
- (57) Girlanda P, Dattola R, Venuto C, Mangiapane R, Nicolosi C, Messina C. Local steroid treatment in idiopathic carpal tunnel syndrome: short- and long-term efficacy. J Neurol 1993;240(3):187-90.
- (58) Gokoglu F, Fndkoglu G, Yorgancoglu ZR, Okumus M, Ceceli E, Kocaoglu S. Evaluation of iontophoresis and local corticosteroid injection in the treatment of carpal tunnel syndrome. Am J Phys Med Rehabil 2005 Feb;84(2):92-6.
- (59) Greenslade JR, Mehta RL, Belward P, Warwick DJ. Dash and Boston questionnaire assessment of carpal tunnel syndrome outcome: what is the responsiveness of an outcome questionnaire? J Hand Surg [Br] 2004 Apr;29(2):159-64.
- (60) Gummesson C, Atroshi I, Ekdahl C. The disabilities of the arm, shoulder and hand (DASH) outcome questionnaire: longitudinal construct validity and measuring self-rated health change after surgery. BMC Musculoskelet Disord 2003 Jun 16;4:11.

- (61) Habib GS, Badarny S, Rawashdeh H. A novel approach of local corticosteroid injection for the treatment of carpal tunnel syndrome. Clin Rheumatol 2006 May;25(3):338-40.
- (62) Hobby JL, Watts C, Elliot D. Validity and responsiveness of the patient evaluation measure as an outcome measure for carpal tunnel syndrome. J Hand Surg [Br] 2005 Aug;30(4):350-4.
- (63) Hui AC, Wong SM, Tang A, Mok V, Hung LK, Wong KS. Long-term outcome of carpal tunnel syndrome after conservative treatment. Int J Clin Pract 2004 Apr;58(4):337-9.
- (64) Hui AC, Wong S, Leung CH, Tong P, Mok V, Poon D, et al. A randomized controlled trial of surgery vs steroid injection for carpal tunnel syndrome. Neurology 2005 Jun 28;64(12):2074-8.
- (65) Imaeda T, Hirata H, Toh S, Nakao Y, Nishida J, Ijichi M, et al. Comparative responsiveness of Japanese versions of the DASH and SF-36 questionnaires and physical measurement to clinical changes after carpal tunnel release. Hand Surg 2006;11(1-2):27-33.
- (66) Jeffrey SL, Belcher HJ. Use of Arnica to relieve pain after carpal-tunnel release surgery. Altern Ther Health Med 2002 Mar;8(2):66-8.
- (67) Jenkinson C, Wright L, Coulter A. Criterion validity and reliability of the SF-36 in a population sample. Qual Life Res 1994 Feb;3(1):7-12.
- (68) Katz JN, Larson MG, Phillips CB, Fossel AH, Liang MH. Comparative measurement sensitivity of short and longer health status instruments. Med Care 1992 Oct;30(10):917-25.
- (69) Keller S, Bann CM, Dodd SL, Schein J, Mendoza TR, Cleeland CS. Validity of the brief pain inventory for use in documenting the outcomes of patients with noncancer pain. Clin J Pain 2004 Sep;20(5):309-18.
- (70) Klein RD, Kotsis SV, Chung KC. Open carpal tunnel release using a 1-centimeter incision: technique and outcomes for 104 patients. Plast Reconstr Surg 2003 Apr 15;111(5):1616-22.
- (71) Kotsis SV, Chung KC. Responsiveness of the Michigan Hand Outcomes Questionnaire and the Disabilities of the Arm, Shoulder and Hand questionnaire in carpal tunnel surgery. J Hand Surg [Am] 2005 Jan;30(1):81-6.
- (72) Kotsis SV, Lau FH, Chung KC. Responsiveness of the Michigan Hand Outcomes Questionnaire and physical measurements in outcome studies of distal radius fracture treatment. J Hand Surg [Am] 2007 Jan;32(1):84-90.

- (73) Leite JC, Jerosch-Herold C, Song F. A systematic review of the psychometric properties of the Boston Carpal Tunnel Questionnaire. BMC Musculoskelet Disord 2006;7:78.
- (74) Lichtman DM, Florio RL, Mack GR. Carpal tunnel release under local anesthesia: evaluation of the outpatient procedure. J Hand Surg [Am] 1979 Nov;4(6):544-6.
- (75) Ly-Pen D, Andreu JL, de BG, Sanchez-Olaso A, Millan I. Surgical decompression versus local steroid injection in carpal tunnel syndrome: a one-year, prospective, randomized, open, controlled clinical trial. Arthritis Rheum 2005 Feb;52(2):612-9.
- (76) MacDermid JC, Richards RS, Donner A, Bellamy N, Roth JH. Responsiveness of the short form-36, disability of the arm, shoulder, and hand questionnaire, patient-rated wrist evaluation, and physical impairment measurements in evaluating recovery after a distal radius fracture. J Hand Surg [Am] 2000 Mar;25(2):330-40.
- (77) Marshall S, Tardif G, Ashworth N. Local corticosteroid injection for carpal tunnel syndrome. Cochrane Database Syst Rev 2002;(4):CD001554.
- (78) Martins RS, Siqueira MG, Simplicio H. Wrist immobilization after carpal tunnel release: A prospective study. Arq Neuro-Psiquiatr 2006;64(3 A).
- (79) Meenan RF, Mason JH, Anderson JJ, Guccione AA, Kazis LE. AIMS2. The content and properties of a revised and expanded Arthritis Impact Measurement Scales Health Status Questionnaire. Arthritis Rheum 1992 Jan;35(1):1-10.
- (80) Menon J. Endoscopic carpal tunnel release: preliminary report. Arthroscopy 1994 Feb;10(1):31-8.
- (81) Michlovitz S, Hun L, Erasala GN, Hengehold DA, Weingand KW. Continuous low-level heat wrap therapy is effective for treating wrist pain. Arch Phys Med Rehabil 2004 Sep;85(9):1409-16.
- (82) Mishra S, Prabhakar S, Lal V, Modi M, Das CP, Khurana D. Efficacy of splinting and oral steroids in the treatment of carpal tunnel syndrome: a prospective randomized clinical and electrophysiological study. Neurol India 2006 Sep;54(3):286-90.
- (83) Mondelli M, Padua L, Reale F, Signorini AM, Romano C. Outcome of surgical release among diabetics with carpal tunnel syndrome. Arch Phys Med Rehabil 2004 Jan;85(1):7-13.
- (84) O'Connor D, Marshall S, Massy-Westropp N. Non-surgical treatment (other than steroid injection) for carpal tunnel syndrome. Cochrane Database Syst Rev 2003;(1):CD003219.

- (85) O'Gradaigh D, Merry P. Corticosteroid injection for the treatment of carpal tunnel syndrome. Ann Rheum Dis 2000 Nov;59(11):918-9.
- (86) Okutsu I, Hamanaka I, Tanabe T, Takatori Y, Ninomiya S. Complete endoscopic carpal tunnel release in long-term haemodialysis patients. J Hand Surg [Br] 1996 Oct;21(5):668-71.
- (87) Osterman AL. The double crush syndrome. Orthop Clin North Am 1988 Jan;19(1):147-55.
- (88) Ozyurekoglu T, McCabe SJ, Goldsmith LJ, LaJoie AS. The minimal clinically important difference of the Carpal Tunnel Syndrome Symptom Severity Scale. J Hand Surg [Am] 2006 May;31(5):733-8.
- (89) Pagnanelli DM, Barrer SJ. Carpal tunnel syndrome: surgical treatment using the Paine retinaculatome. J Neurosurg 1991 Jul;75(1):77-81.
- (90) Paine KW, Polyzoidis KS. Carpal tunnel syndrome. Decompression using the Paine retinaculotome. J Neurosurg 1983 Dec;59(6):1031-6.
- (91) Pinar L, Enhos A, Ada S, Gungor N. Can we use nerve gliding exercises in women with carpal tunnel syndrome? Adv Ther 2005 Sep;22(5):467-75.
- (92) Piravej K, Boonhong J. Effect of ultrasound thermotherapy in mild to moderate carpal tunnel syndrome. J Med Assoc Thai 2004 Sep;87 Suppl 2:S100-S106.
- (93) Premoselli S, Sioli P, Grossi A, Cerri C. Neutral wrist splinting in carpal tunnel syndrome: a 3- and 6-months clinical and neurophysiologic follow-up evaluation of night-only splint therapy. Eura Medicophys 2006 Jun;42(2):121-6.
- (94) Provinciali L, Giattini A, Splendiani G, Logullo F. Usefulness of hand rehabilitation after carpal tunnel surgery. Muscle Nerve 2000 Feb;23(2):211-6.
- (95) Rab M, Grunbeck M, Beck H, Haslik W, Schrogendorfer 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(7):730-6.
- (96a) Reale F, Ginanneschi F, Sicurelli F, Mondelli M. Protocol of outcome evaluation for surgical release of carpal tunnel syndrome. Neurosurgery 2003 Aug;53(2):343-50.
- (96) 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 Oct;28(5):444-9.

- (97) Scholten RJ, Gerritsen AA, Uitdehaag BM, van GD, de Vet HC, Bouter LM. Surgical treatment options for carpal tunnel syndrome. Cochrane Database Syst Rev 2004;(4):CD003905.
- (98) Serra L, Panagiotopoulos K, Bucciero A, Mehrabi FK, Pescatore G, Santangelo M, et al. Endoscopic release in carpal tunnel syndrome: analysis of clinical results in 200 cases. Minim Invasive Neurosurg 2003 Feb;46(1):11-5.
- (99) Sevim S, Dogu O, Camdeviren H, Kaleagasi H, Aral M, Arslan E, et al. Long-term effectiveness of steroid injections and splinting in mild and moderate carpal tunnel syndrome. Neurol Sci 2004 Jun;25(2):48-52.
- (100) Shapiro S. Microsurgical carpal tunnel release. Neurosurgery 1995 Jul;37(1):66-70.
- (101) Siegmeth AW, Hopkinson-Woolley JA. Standard open decompression in carpal tunnel syndrome compared with a modified open technique preserving the superficial skin nerves: a prospective randomized study. J Hand Surg [Am] 2006 Nov;31(9):1483-9.
- (102) Singh A, Gnanalingham K, Casey A, Crockard A. Quality of life assessment using the Short Form-12 (SF-12) questionnaire in patients with cervical spondylotic myelopathy: comparison with SF-36. Spine 2006 Mar 15;31(6):639-43.
- (103) Slattery PG. Endoscopic carpal tunnel release. Use of the modified Chow technique in 215 cases. Med J Aust 1994 Feb 7;160(3):104-7.
- (104) Spies-Dorgelo MN, Terwee CB, Stalman WA, van der Windt DA. Reproducibility and responsiveness of the Symptom Severity Scale and the hand and finger function subscale of the Dutch arthritis impact measurement scales (Dutch-AIMS2-HFF) in primary care patients with wrist or hand problems. Health Qual Life Outcomes 2006;4:87.
- (105) Straub TA. Endoscopic carpal tunnel release: a prospective analysis of factors associated with unsatisfactory results. Arthroscopy 1999 Apr;15(3):269-74.
- (106) Tan G, Jensen MP, Thornby JI, Shanti BF. Validation of the Brief Pain Inventory for chronic nonmalignant pain. J Pain 2004 Mar;5(2):133-7.
- (107) 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 Oct;114(5):1137-46.
- (108) Ucan H, Yagci I, Yilmaz L, Yagmurlu F, Keskin D, Bodur H. Comparison of splinting, splinting plus local steroid injection and open carpal tunnel release outcomes in idiopathic carpal tunnel syndrome. Rheumatol Int 2006 Nov;27(1):45-51.

- (109) Upton AR, McComas AJ. The double crush in nerve entrapment syndromes. Lancet 1973 Aug 18;2(7825):359-62.
- (110) Verdugo RJ, Salinas RS, Castillo J, Cea JG. Surgical versus non-surgical treatment for carpal tunnel syndrome. Cochrane Database Syst Rev 2003;(3):CD001552.
- (111) Waegeneers S, Haentjens P, Wylock P. Operative treatment of carpal tunnel syndrome. Acta Orthop Belg 1993;59(4):367-70.
- (112) Wah JW, Wang MK, Ping CL. Construct validity of the Chinese version of the Patient-rated Wrist Evaluation Questionnaire (PRWE-Hong Kong Version). J Hand Ther 2006 Jan;19(1):18-26, quiz.
- (113) Wilgis EF, Burke FD, Dubin NH, Sinha S, Bradley MJ. A prospective assessment of carpal tunnel surgery with respect to age. J Hand Surg [Br] 2006 Aug;31(4):401-6.
- (114) Williams VS, Smith MY, Fehnel SE. The validity and utility of the BPI interference measures for evaluating the impact of osteoarthritic pain. J Pain Symptom Manage 2006 Jan;31(1):48-57.
- (115) 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 Aug;85(6):863-8.
- (116) Wong SM, Hui AC, Lo SK, Chiu JH, Poon WF, Wong L. Single vs. two steroid injections for carpal tunnel syndrome: a randomised clinical trial. Int J Clin Pract 2005 Dec;59(12):1417-21.

REFERENCES TO EXCLUDED ARTICLES

- (117) Abasolo L, Carmona L, Hernandez-Garcia C, Lajas C, Loza E, Blanco M, et al. Musculoskeletal work disability for clinicians: time course and effectiveness of a specialized intervention program by diagnosis. Arthritis Rheum 2007 Mar 15;57(2):335-42.
- (118) Abdullah AF, Wolber PH, Ditto EW, III. Sequelae of carpal tunnel surgery: rationale for the design of a surgical approach. Neurosurgery 1995 Nov;37(5):931-5.
- (119) Adams ML, Franklin GM, Barnhart S. Outcome of carpal tunnel surgery in Washington State workers' compensation. Am J Ind Med 1994 Apr;25(4):527-36.
- (120) Agarwal V, Singh R, Sachdev A, Wiclaff, Shekhar S, Goel D. A prospective study of the long-term efficacy of local methyl prednisolone acetate injection in the management of mild carpal tunnel syndrome. Rheumatology (Oxford) 2005 May;44(5):647-50.

- (121) 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 Nov;17(6):987-95.
- (122) Ahcan U, Arnez ZM, Bajrovic F, Zorman P. Surgical technique to reduce scar discomfort after carpal tunnel surgery. J Hand Surg [Am] 2002 Sep;27(5):821-7.
- (123) Akalin E, El O, Peker O, Senocak O, Tamci S, Gulbahar S, et al. Treatment of carpal tunnel syndrome with nerve and tendon gliding exercises. Am J Phys Med Rehabil 2002 Feb;81(2):108-13.
- (124) Alayurt S, Memis D, Pamukcu Z. The addition of sufentanil, tramadol or clonidine to lignocaine for intravenous regional anaesthesia. Anaesth Intensive Care 2004 Feb;32(1):22-7.
- (125) Altissimi M, Mancini GB. Surgical release of the median nerve under local anaesthesia for carpal tunnel syndrome. J Hand Surg [Br] 1988 Nov;13(4):395-6.
- (126) Andreu JL, Ly-Pen D. A randomized controlled trial of surgery vs steroid injection for carpal tunnel syndrome. Neurology 2006 Mar 28;66(6):955-6.
- (127) Anonymous. Steroid injection equivalent to surgery for carpal tunnel syndrome. J Fam Pract 2005;54(5):401.
- (128) Ariyan S, Watson HK. The palmar approach for the visualization and release of the carpal tunnel. An analysis of 429 cases. Plast Reconstr Surg 1977 Oct;60(4):539-47.
- (129) Armitage M, Shepherd S. A new professional in the healthcare workforce: role, training, assessment and regulation. Clin Med 2005 Jul;5(4):311-4.
- (130) Atherton WG, Faraj AA, Riddick AC, Davis TR. Follow-up after carpal tunnel decompression general practitioner surgery or hand clinic? A randomized prospective study. J Hand Surg [Br] 1999 Jun;24(3):296-7.
- (131) Atroshi I, Johnsson R, Ornstein E. Patient satisfaction and return to work after endoscopic carpal tunnel surgery. J Hand Surg [Am] 1998 Jan;23(1):58-65.
- (132) Baba H, Maezawa Y, Uchida K, Furusawa N, Wada M, Imura S, et al. Cervical myeloradiculopathy with entrapment neuropathy: a study based on the double-crush concept. Spinal Cord 1998 Jun;36(6):399-404.
- (133) Banta CA. A prospective, nonrandomized study of iontophoresis, wrist splinting, and antiinflammatory medication in the treatment of early-mild carpal tunnel syndrome. J Occup Med 1994 Feb;36(2):166-8.

- (134) Benedetti VR, Sennwald G. [Agee endoscopic decompression of the median nerve: prospective study with comparison to open decompression]. Handchir Mikrochir Plast Chir 1996 May;28(3):151-5.
- (135) Bessette L, Keller RB, Liang MH, Simmons BP, Fossel AH, Katz JN. Patients' preferences and their relationship with satisfaction following carpal tunnel release. J Hand Surg [Am] 1997 Jul;22(4):613-20.
- (136) Bhatia R, Field J, Grote J, Huma H. Does splintage help pain after carpal tunnel release? J Hand Surg [Br] 2000 Apr;25(2):150.
- (137) Bigat Z, Karsli B, Boztug N, Cete N, Ertok E. Comparison of the effect of low-dose ropivacaine and lidocaine in intravenous regional anaesthesia: A randomised, double-blind clinical study. Clin Drug Invest 2005;25(3):209-14.
- (138) Bigat Z, Boztug N, Hadimioglu N, Cete N, Coskunfirat N, Ertok E. Does dexamethasone improve the quality of intravenous regional anesthesia and analgesia? A randomized, controlled clinical study. Anesth Analg 2006 Feb;102(2):605-9.
- (139) Birkbeck MQ, Beer TC. Occupation in relation to the carpal tunnel syndrome. Rheumatol Rehabil 1975 Nov;14(4):218-21.
- (140) Bitar G, Alexandrides J, Missirian R, Sotereanos D, Nystrom A. Carpal tunnel release in the United States and Sweden: reimbursement patterns, cost for treatment, and return to work. Plast Reconstr Surg 2002 Apr 15;109(5):1574-8.
- (141) Bland JD. Do nerve conduction studies predict the outcome of carpal tunnel decompression? Muscle Nerve 2001 Jul;24(7):935-40.
- (142) Bonfiglioli R, Mattioli S, Fiorentini C, Graziosi F, Curti S, Violante FS. Relationship between repetitive work and the prevalence of carpal tunnel syndrome in part-time and full-time female supermarket cashiers: a quasi-experimental study. Int Arch Occup Environ Health 2007 Jan;80(3):248-53.
- (143) Bonzani PJ, Millender L, Keelan B, Mangieri MG. Factors prolonging disability in work-related cumulative trauma disorders. J Hand Surg [Am] 1997 Jan;22(1):30-4.
- (144) Bovenzi M, Zadini A, Franzinelli A, Borgogni F. Occupational musculoskeletal disorders in the neck and upper limbs of forestry workers exposed to hand-arm vibration. Ergonomics 1991 May;34(5):547-62.
- (145) Boyd KU, Gan BS, Ross DC, Richards RS, Roth JH, MacDermid JC. Outcomes in carpal tunnel syndrome: symptom severity, conservative management and progression to surgery. Clin Invest Med 2005 Oct;28(5):254-60.

- (146) Braga Silva J, Fontes Neto P, Foucher G, Fridman M. Postoperative strength after surgical release of the carpal tunnel: A randomized prospective study. REV BRAS ORTOP 1996;31(4):355-7.
- (147) Braun RM, Doehr S, Mosqueda T, Garcia A. The effect of legal representation on functional recovery of the hand in injured workers following carpal tunnel release. J Hand Surg [Am] 1999 Jan;24(1):53-8.
- (148) Breuer B, Sperber K, Wallenstein S, Kiprovski K, Calapa A, Snow B, et al. Clinically significant placebo analgesic response in a pilot trial of botulinum B in patients with hand pain and carpal tunnel syndrome. Pain Med 2006 Jan;7(1):16-24.
- (149) Brown MG, Keyser B, Rothenberg ES. Endoscopic carpal tunnel release. J Hand Surg [Am] 1992 Nov;17(6):1009-11.
- (150) Brown MG, Rothenberg ES, Keyser B, Woloszyn TT, Wolford A. Results of 1236 endoscopic carpal tunnel release procedures using the Brown technique. Contemp Orthop 1993 Sep;27(3):251-8.
- (151) Brown RA, Gelberman RH, Seiler JG, III, 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 Sep;75(9):1265-75.
- (152) Burke FD, Bradley MJ, Sinha S, Wilgis EF, Dubin NH. Primary care management of patients with carpal tunnel syndrome referred to surgeons: are non-operative interventions effectively utilised? Postgrad Med J 2007 Jul;83(981):498-501.
- (153) Carr AJ, Higginson IJ. Are quality of life measures patient centred? BMJ 2001 Jun 2;322(7298):1357-60.
- (154) Carroll RE, Hurst LC. The relationship of thoracic outlet syndrome and carpal tunnel syndrome. Clin Orthop Relat Res 1982 Apr;(164):149-53.
- (155) Carter R, Aspy CB, Mold J. The effectiveness of magnet therapy for treatment of wrist pain attributed to carpal tunnel syndrome. J Fam Pract 2002 Jan;51(1):38-40.
- (156) Cavallo AV, Slattery PG, Barton RJ. Endoscopic carpal tunnel release and congenital anomalies of the median nerve. Hand Surg 2003 Dec;8(2):265-70.
- (157) Celiker R, Arslan S, Inanici F. Corticosteroid injection vs. nonsteroidal antiinflammatory drug and splinting in carpal tunnel syndrome. Am J Phys Med Rehabil 2002 Mar;81(3):182-6.
- (158) Chaise F, Bellemere P, Fril JP, Gaisne E, Poirier P, Menadi A. Return-to-work interval and surgery for carpal tunnel syndrome. Results of a prospective series of 233 patients. J Hand Surg [Br] 2004 Dec;29(6):568-70.

- (159) Chan L, Turner JA, Comstock BA, Levenson LM, Hollingworth W, Heagerty PJ, et al. The relationship between electrodiagnostic findings and patient symptoms and function in carpal tunnel syndrome. Arch Phys Med Rehabil 2007 Jan;88(1):19-24.
- (160) Chang MH, Ger LP, Hsieh PF, Huang SY. A randomised clinical trial of oral steroids in the treatment of carpal tunnel syndrome: a long term follow up. J Neurol Neurosurg Psychiatry 2002 Dec;73(6):710-4.
- (161) Changulani M, Okonkwo U, Keswani T, Kalairajah Y. Outcome evaluation measures for wrist and hand which one to choose? Int Orthop 2007 May 30.
- (162) Chen HT, Chen HC, Wei FC. Endoscopic carpal tunnel release. Changgeng Yi Xue Za Zhi 1999 Sep;22(3):386-91.
- (163) Chertow GM, Trimbur T, Karlson EW, Lazarus JM, Kay J. Performance characteristics of a dialysis-related amyloidosis questionnaire. J Am Soc Nephrol 1996 Aug;7(8):1235-40.
- (164) Chow JC. Endoscopic release of the carpal ligament for carpal tunnel syndrome: 22-month clinical result. Arthroscopy 1990;6(4):288-96.
- (165) Chow JC, Hantes ME. Endoscopic carpal tunnel release: thirteen years' experience with the Chow technique. J Hand Surg [Am] 2002 Nov;27(6):1011-8.
- (166) Concannon MJ, Gainor B, Petroski GF, Puckett CL. The predictive value of electrodiagnostic studies in carpal tunnel syndrome. Plast Reconstr Surg 1997 Nov;100(6):1452-8.
- (167) Concannon MJ, Brownfield ML, Puckett CL. The incidence of recurrence after endoscopic carpal tunnel release. Plast Reconstr Surg 2000 Apr;105(5):1662-5.
- (168) Cosgrove JL, Chase PM, Mast NJ, Reeves R. Carpal tunnel syndrome in railroad workers. Am J Phys Med Rehabil 2002 Feb;81(2):101-7.
- (169) Dakowicz A, Latosiewicz R. The value of iontophoresis combined with ultrasound in patients with the carpal tunnel syndrome. Rocz Akad Med Bialymst 2005;50 Suppl 1:196-8.
- (170) Das SK, Brown HG. In search of complications in carpal tunnel decompression. Hand 1976 Oct;8(3):243-9.
- (171) Davis AM, Beaton DE, Hudak P, Amadio P, Bombardier C, Cole D, et al. Measuring disability of the upper extremity: a rationale supporting the use of a regional outcome measure. J Hand Ther 1999 Oct;12(4):269-74.

- (172) Davis L, Wellman H, Punnett L. Surveillance of work-related carpal tunnel syndrome in Massachusetts, 1992-1997: a report from the Massachusetts Sentinel Event Notification System for Occupational Risks (SENSOR). Am J Ind Med 2001 Jan;39(1):58-71.
- (173) Davis PT, Hulbert JR, Kassak KM, Meyer JJ. Comparative efficacy of conservative medical and chiropractic treatments for carpal tunnel syndrome: a randomized clinical trail. J Manipulative Physiol Ther 1998 Jun;21(5):317-26.
- (174) Dawson WJ, Elenz DR, Winchester DP, Feldman JL. Elective hand surgery in the breast cancer patient with prior ipsilateral axillary dissection. Ann Surg Oncol 1995 Mar;2(2):132-7.
- (175) Deyo RA, Centor RM. Assessing the responsiveness of functional scales to clinical change: an analogy to diagnostic test performance. J Chronic Dis 1986;39(11):897-906.
- (176) Drapela J, Syrovy J, Kulakovska M. Carpal tunnel syndrome. Revision of the thenar motor branch? Acta Chir Plast 1997;39(3):78-81.
- (177) Duchateau JA, Moermans JP. Carpal tunnel syndrome: postsurgical course of symptoms. Ann Chir Main 1984;3(3):227-31.
- (178) Dumontier C, Sokolow C, Leclercq C, Chauvin P. Early results of conventional versus two-portal endoscopic carpal tunnel release. A prospective study. J Hand Surg [Br] 1995 Oct;20(5):658-62.
- (179) Eason SY, Belsole RJ, Greene TL. Carpal tunnel release: analysis of suboptimal results. J Hand Surg [Br] 1985 Oct;10(3):365-9.
- (180) Ebenbichler GR, 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 Mar 7;316(7133):731-5.
- (181) Edgell SE, McCabe SJ, Breidenbach WC, LaJoie AS, Abell TD. Predicting the outcome of carpal tunnel release. J Hand Surg [Am] 2003 Mar;28(2):255-61.
- (182) English CJ, Maclaren WM, Court-Brown, Hughes SP, Porter RW, Wallace WA, et al. Relations between upper limb soft tissue disorders and repetitive movements at work. Am J Ind Med 1995 Jan;27(1):75-90.
- (183) Erdmann MW. Endoscopic carpal tunnel decompression. J Hand Surg [Br] 1994 Feb;19(1):5-13.
- (184) Erel E, Pleasance PI, Ahmed O, Hart NB. Absorbable versus non-absorbable suture in carpal tunnel decompression. J Hand Surg [Br] 2001 Apr;26(2):157-8.

- (185) Eskandari MM, Ozge A, Oztuna V, Colak M, Kanik A, Kuyurtar F. Effect of patient age and symptom duration on subjective and objective outcomes of carpal tunnel surgery. Orthopedics 2005 Jun;28(6):600-2.
- (186) Eversmann W. Reduction of cumulative trauma disorders by a comprehensive ergonomic program in a major commercial bakery. ASSH 1990;9:1-8.
- (187) Ferdinand RD, MacLean JG. Endoscopic versus open carpal tunnel release in bilateral carpal tunnel syndrome. A prospective, randomised, blinded assessment. J Bone Joint Surg Br 2002 Apr;84(3):375-9.
- (188) Feuerstein M, Miller VL, Burrell LM, Berger R. Occupational upper extremity disorders in the federal workforce. Prevalence, health care expenditures, and patterns of work disability. J Occup Environ Med 1998 Jun;40(6):546-55.
- (189) Feuerstein M, Burrell LM, Miller VI, Lincoln A, Huang GD, Berger R. Clinical management of carpal tunnel syndrome: a 12-year review of outcomes. Am J Ind Med 1999 Mar;35(3):232-45.
- (190) Fish DR, Morris-Allen DM. Musculoskeletal disorders in dentists. N Y State Dent J 1998 Apr;64(4):44-8.
- (191) Fissette J, Onkelinx A. Treatment of carpal tunnel syndrome. Comparative study with and without epineurolysis. Hand 1979 Jun;11(2):206-10.
- (192) Flak M, Durmala J, Czernicki K, Dobosiewicz K. Double crush syndrome evaluation in the median nerve in clinical, radiological and electrophysiological examination. Stud Health Technol Inform 2006;123:435-41.
- (193) Flaschka G, Eder H, Mullegger G, Gindl HK. Follow-up results of surgery for carpal tunnel syndrome in local anesthesia. Zentralbl Neurochir 1991;52(3):123-5.
- (194) Foucher G, Buch N, Van OL, Gautherie M, Jesel M. [Carpal tunnel syndrome. Can it still be a controversial topic?]. Chirurgie 1993;119(1-2):80-4.
- (195) Franklin GM, Fulton-Kehoe D. Outcomes research in Washington state workers' compensation. Am J Ind Med 1996 Jun;29(6):642-8.
- (196) Garfinkel MS, Singhal A, Katz WA, Allan DA, Reshetar R, Schumacher HR, Jr. Yoga-based intervention for carpal tunnel syndrome: a randomized trial. JAMA 1998 Nov 11;280(18):1601-3.
- (197) GARLAND H, LANGWORTH EP, TAVERNER D, CLARK JM. SURGICAL TREATMENT FOR THE CARPAL TUNNEL SYNDROME. Lancet 1964 May 23;1:1129-30.

- (198) Gell N, Werner RA, Franzblau A, Ulin SS, Armstrong TJ. A longitudinal study of industrial and clerical workers: incidence of carpal tunnel syndrome and assessment of risk factors. J Occup Rehabil 2005 Mar;15(1):47-55.
- (199) Gerr F, Letz R, Landrigan PJ. Upper-extremity musculoskeletal disorders of occupational origin. Annu Rev Public Health 1991;12:543-66.
- (200) Gerr F, Marcus M, Monteilh C, Hannan L, Ortiz D, Kleinbaum D. A randomised controlled trial of postural interventions for prevention of musculoskeletal symptoms among computer users. Occup Environ Med 2005 Jul;62(7):478-87.
- (201) Gerritsen AA, Scholten RJ, Assendelft WJ, Kuiper H, de Vet HC, Bouter LM. Splinting or surgery for carpal tunnel syndrome? Design of a randomized controlled trial [ISRCTN18853827]. BMC Neurol 2001 Dec 18;1:8.
- (202) 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 Sep 11;288(10):1245-51.
- (203) Gerritsen AA, de Krom MC, Struijs MA, Scholten RJ, de Vet HC, Bouter LM. Conservative treatment options for carpal tunnel syndrome: a systematic review of randomised controlled trials. J Neurol 2002 Mar;249(3):272-80.
- (204) Gerritsen AA, Korthals-de B, I, Laboyrie PM, de Vet HC, Scholten RJ, Bouter LM. Splinting for carpal tunnel syndrome: prognostic indicators of success. J Neurol Neurosurg Psychiatry 2003 Sep;74(9):1342-4.
- (205) Giele H. Evidence-based treatment of carpal tunnel syndrome. Curr Orthop 2001;15(4):249-55.
- (206) Giersiepen K, Eberle A, Pohlabeln H. Gender differences in carpal tunnel syndrome? occupational and non-occupational risk factors in a population-based case-control study. Ann Epidemiol 2000 Oct 1;10(7):481.
- (207) Gimeno D, Amick BC, III, Habeck RV, Ossmann J, Katz JN. The role of job strain on return to work after carpal tunnel surgery. Occup Environ Med 2005 Nov;62(11):778-85.
- (208) Goitz RJ, Steichen JB. Microvascular omental transfer for the treatment of severe recurrent median neuritis of the wrist: a long-term follow-up. Plast Reconstr Surg 2005 Jan;115(1):163-71.
- (209) Graham BA. Two weeks of prednisolone was as effective as four weeks in improving carpal tunnel syndrome symptoms. J Bone Joint Surg Am 2003 Aug;85-A(8):1624.

- (210) Graham RG, Hudson DA, Solomons M, Singer M. A prospective study to assess the outcome of steroid injections and wrist splinting for the treatment of carpal tunnel syndrome. Plast Reconstr Surg 2004 Feb;113(2):550-6.
- (211) Greening J. Workshop: clinical implications for clinicians treating patients with non-specific arm pain, whiplash and carpal tunnel syndrome. Man Ther 2006 Aug;11(3):171-2.
- (212) Gunetti R, Bonicalzi V, Riolo C, Pagni CA. Peri- and postoperative pain valutation in carpal tunnel release of median nerve compression. J Neurosurg Sci 2000 Jun;44(2):85-8.
- (213) Guyatt G, Walter S, Norman G. Measuring change over time: assessing the usefulness of evaluative instruments. J Chronic Dis 1987;40(2):171-8.
- (214) Guyatt GH, Deyo RA, Charlson M, Levine MN, Mitchell A. Responsiveness and validity in health status measurement: a clarification. J Clin Epidemiol 1989;42(5):403-8.
- (215) Hagebeuk EE, de Weerd AW. Clinical and electrophysiological follow-up after local steroid injection in the carpal tunnel syndrome. Clin Neurophysiol 2004 Jun;115(6):1464-8.
- (216) Hanssen AD, Amadio PC, DeSilva SP, Ilstrup DM. Deep postoperative wound infection after carpal tunnel release. J Hand Surg [Am] 1989 Sep;14(5):869-73.
- (217) Helm RH, Vaziri S. Evaluation of carpal tunnel release using the Knifelight instrument. J Hand Surg [Br] 2003 Jun;28(3):251-4.
- (218) Herbert R, Gerr F, Dropkin J. Clinical evaluation and management of work-related carpal tunnel syndrome. Am J Ind Med 2000 Jan;37(1):62-74.
- (219) Heuser A, Kourtev H, Winter S, Fensterheim D, Burdea G, Hentz V, et al. Telerehabilitation using the Rutgers Master II glove following carpal tunnel release surgery: proof-of-concept. IEEE Trans Neural Syst Rehabil Eng 2007 Mar;15(1):43-9.
- (220) Heybeli N, Kutluhan S, Demirci S, Kerman M, Mumcu EF. Assessment of outcome of carpal tunnel syndrome: a comparison of electrophysiological findings and a self-administered Boston questionnaire. J Hand Surg [Br] 2002 Jun;27(3):259-64.
- (221) Higgs PE, Edwards D, Martin DS, Weeks PM. Carpal tunnel surgery outcomes in workers: effect of workers' compensation status. J Hand Surg [Am] 1995 May;20(3):354-60.

- (222) Hochberg J. A randomized prospective study to assess the efficacy of two cold-therapy treatments following carpal tunnel release. J Hand Ther 2001 Jul;14(3):208-15.
- (223) Hoefnagels WA, van Kleef JG, Mastenbroek GG, de Blok JA, Breukelman AJ, de Krom MC. [Surgical treatment of carpal tunnel syndrome: endoscopic or classical (open)? A prospective randomized trial]. Ned Tijdschr Geneeskd 1997 May 3;141(18):878-82.
- (224) Howard FM. Compression neuropathies in the anterior forearm. Hand Clin 1986 Nov;2(4):737-45.
- (225) Hudak PL, Amadio PC, Bombardier C. Development of an upper extremity outcome measure: the DASH (disabilities of the arm, shoulder and hand) [corrected]. The Upper Extremity Collaborative Group (UECG). Am J Ind Med 1996 Jun;29(6):602-8.
- (226) Hui AC, Wong SM, Wong KS, Li E, Kay R, Yung P, et al. Oral steroid in the treatment of carpal tunnel syndrome. Ann Rheum Dis 2001 Aug;60(8):813-4.
- (227) Huracek J, Heising T, Wanner M, Troeger H. Recovery after carpal tunnel syndrome operation: the influence of the opposite hand, if operated on in the same session. Arch Orthop Trauma Surg 2001 Jul;121(7):368-70.
- (228) Hurst LC, Weissberg D, Carroll RE. The relationship of the double crush to carpal tunnel syndrome (an analysis of 1,000 cases of carpal tunnel syndrome). J Hand Surg [Br] 1985 Jun;10(2):202-4.
- (229) Imaeda T, Toh S, Wada T, Uchiyama S, Okinaga S, Kusunose K, et al. Validation of the Japanese Society for Surgery of the Hand Version of the Quick Disability of the Arm, Shoulder, and Hand (QuickDASH-JSSH) questionnaire. J Orthop Sci 2006 May;11(3):248-53.
- (230) Irvine J, Chong SL, Amirjani N, Chan KM. Double-blind randomized controlled trial of low-level laser therapy in carpal tunnel syndrome. Muscle Nerve 2004 Aug;30(2):182-7.
- (231) 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 Apr;21(2):202-4.
- (232) Jensen MP, Gammaitoni AR, Olaleye DO, Oleka N, Nalamachu SR, Galer BS. The pain quality assessment scale: assessment of pain quality in carpal tunnel syndrome. J Pain 2006 Nov;7(11):823-32.
- (233) Jerosch-Herold C, Leite JC, Song F. A systematic review of outcomes assessed in randomized controlled trials of surgical interventions for carpal tunnel syndrome using

- the International Classification of Functioning, Disability and Health (ICF) as a reference tool. BMC Musculoskelet Disord 2006;7:96.
- (234) Jimenez DF, Gibbs SR, Clapper AT. Endoscopic treatment of carpal tunnel syndrome: a critical review. J Neurosurg 1998 May;88(5):817-26.
- (235) Kane RL, Bershadsky B, Lin WC, Rockwood T, Wood K. Efforts to standardize the reporting of pain. J Clin Epidemiol 2002 Feb;55(2):105-10.
- (236) Kane RL, Bershadsky B, Rockwood T, Saleh K, Islam NC. Visual Analog Scale pain reporting was standardized. J Clin Epidemiol 2005 Jun;58(6):618-23.
- (237) Kastrup O. Steroid injection or surgery for the treatment of carpal tunnel syndrome? Comment. MMW-Fortschr Med 2007;149(1-2):26.
- (238) Katz JN, Gelberman RH, Wright EA, Abrahamsson SO, Lew RA. A preliminary scoring system for assessing the outcome of carpal tunnel release. J Hand Surg [Am] 1994 Jul;19(4):531-8.
- (239) Katz JN, Keller RB, Simmons BP, Rogers WD, Bessette L, Fossel AH, et al. Maine Carpal Tunnel Study: outcomes of operative and nonoperative therapy for carpal tunnel syndrome in a community-based cohort. J Hand Surg [Am] 1998 Jul;23(4):697-710.
- (240) Katz JN, Losina E, Amick BC, III, Fossel AH, Bessette L, Keller RB. Predictors of outcomes of carpal tunnel release. Arthritis Rheum 2001 May;44(5):1184-93.
- (241) Katz JN, Amick BC, III, Keller R, Fossel AH, Ossman J, Soucie V, et al. Determinants of work absence following surgery for carpal tunnel syndrome. Am J Ind Med 2005 Feb;47(2):120-30.
- (242) Kearns J, Gresch EE, Weichel CY, Eby P, Pallapothu SR. Pre- and post-employment median nerve latency in pork processing employees. J Occup Environ Med 2000 Jan;42(1):96-100.
- (243) Keogh JP, Gucer PW, Gordon JL, Nuwayhid I. Patterns and predictors of employer risk-reduction activities (ERRAs) in response to a work-related upper extremity cumulative trauma disorder (UECTD): reports from workers' compensation claimants. Am J Ind Med 2000 Nov;38(5):489-97.
- (244) Ketchum LD. A comparison of flexor tenosynovectomy, open carpal tunnel release, and open carpal tunnel release with flexor tenosynovectomy in the treatment of carpal tunnel syndrome. Plast Reconstr Surg 2004 Jun;113(7):2020-9.

- (245) Kharwadkar N, Naique S, Molitor PJ. Prospective randomized trial comparing absorbable and non-absorbable sutures in open carpal tunnel release. J Hand Surg [Br] 2005 Feb;30(1):92-5.
- (246) Korthals-de B, I, Gerritsen AA, van Tulder MW, Rutten-van Molken MP, Ader HJ, de Vet HC, et al. Surgery is more cost-effective than splinting for carpal tunnel syndrome in the Netherlands: results of an economic evaluation alongside a randomized controlled trial. BMC Musculoskelet Disord 2006;7:86.
- (247) Kwon HK, Hwang M, Yoon DW. Frequency and severity of carpal tunnel syndrome according to level of cervical radiculopathy: double crush syndrome? Clin Neurophysiol 2006 Jun;117(6):1256-9.
- (248) Laso Guzman FJ, Gonzalez-Buitrago JM, de AF, Mateos F, Moyano JC, Lopez-Alburquerque T. Carpal tunnel syndrome and vitamin B6. Klin Wochenschr 1989 Jan 4;67(1):38-41.
- (249) Lawrence TM, Desai VV. Topical anaesthesia to reduce pain associated with carpal tunnel surgery. J Hand Surg [Br] 2002 Oct;27(5):462-4.
- (250) Leclerc A, Landre MF, Chastang JF, Niedhammer I, Roquelaure Y. Upper-limb disorders in repetitive work. Scand J Work Environ Health 2001 Aug;27(4):268-78.
- (251) Lee H, Jackson TA. Carpal tunnel release through a limited skin incision under direct visualization using a new instrument, the carposcope. Plast Reconstr Surg 1996 Aug;98(2):313-9.
- (252) Li S, Liu L, Miyazaki M, Warren S. Effectiveness of splinting for work-related carpal tunnel syndrome: A three-month follow-up study. Technol Disabil 1999;11(1-2):51-64.
- (253) Lincoln AE, Feuerstein M, Shaw WS, Miller VI. Impact of case manager training on worksite accommodations in workers' compensation claimants with upper extremity disorders. J Occup Environ Med 2002 Mar;44(3):237-45.
- (254) Lorgelly PK, Dias JJ, Bradley MJ, Burke FD. Carpal tunnel syndrome, the search for a cost-effective surgical intervention: a randomised controlled trial. Ann R Coll Surg Engl 2005 Jan;87(1):36-40.
- (255) Macaire P, Choquet O, Jochum D, Travers V, Capdevila X. Nerve blocks at the wrist for carpal tunnel release revisited: the use of sensory-nerve and motor-nerve stimulation techniques. Reg Anesth Pain Med 2005 Nov;30(6):536-40.
- (256) MacDermid J. A hand brace improve symptoms and function in carpal tunnel syndrome. Aust J Physiother 2002;48(2):134.

- (257) MacDermid JC, Turgeon T, Richards RS, Beadle M, Roth JH. Patient rating of wrist pain and disability: a reliable and valid measurement tool. J Orthop Trauma 1998 Nov;12(8):577-86.
- (258) Mackenzie DJ, Hainer R, Wheatley MJ. Early recovery after endoscopic vs. short-incision open carpal tunnel release. Ann Plast Surg 2000 Jun;44(6):601-4.
- (259) Makowiec RL, Nagle DJ, Chow JC. Outcome of first-time endoscopic carpal tunnel release in a teaching environment. Arthroscopy 2002 Jan;18(1):27-31.
- (260) Manente G, Torrieri F, Di BF, Staniscia T, Romano F, Uncini A. An innovative hand brace for carpal tunnel syndrome: a randomized controlled trial. Muscle Nerve 2001 Aug;24(8):1020-5.
- (261) Martin BI, Levenson LM, Hollingworth W, Kliot M, Heagerty PJ, Turner JA, et al. Randomized clinical trial of surgery versus conservative therapy for carpal tunnel syndrome [ISRCTN84286481]. BMC Musculoskelet Disord 2005;6:2.
- (262) Massey EW, Riley TL, Pleet AB. Coexistent carpal tunnel syndrome and cervical radiculopathy (double crush syndrome). South Med J 1981 Aug;74(8):957-9.
- (263) McDonough JW, Gruenloh TJ. A comparison of endoscopic and open carpal tunnel release. Wis Med J 1993 Dec;92(12):675-7.
- (264) McHorney CA, Ware JE, Jr., Raczek AE. The MOS 36-Item Short-Form Health Survey (SF-36): II. Psychometric and clinical tests of validity in measuring physical and mental health constructs. Med Care 1993 Mar;31(3):247-63.
- (265) McNally SA, Hales PF. Results of 1245 endoscopic carpal tunnel decompressions. Hand Surg 2003 Jul;8(1):111-6.
- (266) Menon J. Endoscopic carpal tunnel release: a single-portal technique. Contemp Orthop 1993 Feb;26(2):109-16.
- (267) Menon J, Etter C. Endoscopic carpal tunnel release--current status. J Hand Ther 1993 Apr;6(2):139-44.
- (268) Menovsky T, Bartels RH, van Lindert EL, Grotenhuis JA. Skin closure in carpal tunnel surgery: a prospective comparative study between nylon, polyglactin 910 and stainless steel sutures. Hand Surg 2004 Jul;9(1):35-8.
- (269) Mirza MA, King ET, Jr., Tanveer S. Palmar uniportal extrabursal endoscopic carpal tunnel release. Arthroscopy 1995 Feb;11(1):82-90.

- (270) Mondelli M, Reale F, Sicurelli F, Padua L. Relationship between the self-administered Boston questionnaire and electrophysiological findings in follow-up of surgically-treated carpal tunnel syndrome. J Hand Surg [Br] 2000 Apr;25(2):128-34.
- (271) Mondelli M, Aprile I, Ballerini M, Ginanneschi F, Reale F, Romano C, et al. Sex differences in carpal tunnel syndrome: comparison of surgical and non-surgical populations. Eur J Neurol 2005 Dec;12(12):976-83.
- (272) Mondelli M, Padua L, Giannini F, Bibbo G, Aprile I, Rossi S. A self-administered questionnaire of ulnar neuropathy at the elbow. Neurol Sci 2006 Dec;27(6):402-11.
- (273) Monsivais JJ, Bucher PA, Monsivais DB. Nonsurgically treated carpal tunnel syndrome in the manual worker. Plast Reconstr Surg 1994 Oct;94(5):695-8.
- (274) Morgan G, Wilbourn AJ. Cervical radiculopathy and coexisting distal entrapment neuropathies: double-crush syndromes? Neurology 1998 Jan;50(1):78-83.
- (275) Morgenstern H, Kelsh M, Kraus J, Margolis W. A cross-sectional study of hand/wrist symptoms in female grocery checkers. Am J Ind Med 1991;20(2):209-18.
- (276) Morse LH. Repetitive motion musculoskeletal problems in the microelectronics industry. Occup Med 1986 Jan;1(1):167-74.
- (277) Naeser MA, Hahn KA, Lieberman BE, Branco KF. Carpal tunnel syndrome pain treated with low-level laser and microamperes transcutaneous electric nerve stimulation: A controlled study. Arch Phys Med Rehabil 2002 Jul;83(7):978-88.
- (278) Nalamachu S, Crockett RS, Gammaitoni AR, Gould EM. A comparison of the lidocaine patch 5% vs naproxen 500 mg twice daily for the relief of pain associated with carpal tunnel syndrome: a 6-week, randomized, parallel-group study. MedGenMed 2006;8(3):33.
- (279) Nalamachu S, Crockett RS, Mathur D. Lidocaine patch 5 for carpal tunnel syndrome: how it compares with injections: a pilot study. J Fam Pract 2006 Mar;55(3):209-14.
- (280) Nathan PA, Meadows KD, Keniston RC. Rehabilitation of carpal tunnel surgery patients using a short surgical incision and an early program of physical therapy. J Hand Surg [Am] 1993 Nov;18(6):1044-50.
- (281) Nathan PA, Meadows KD, Istvan JA. Predictors of carpal tunnel syndrome: an 11-year study of industrial workers. J Hand Surg [Am] 2002 Jul;27(4):644-51.
- (282) Nemoto K, Matsumoto N, Tazaki K, Horiuchi Y, Uchinishi K, Mori Y. An experimental study on the "double crush" hypothesis. J Hand Surg [Am] 1987 Jul;12(4):552-9.

- (283) Netscher D, Steadman AK, Thornby J, Cohen V. Temporal changes in grip and pinch strength after open carpal tunnel release and the effect of ligament reconstruction. J Hand Surg [Am] 1998 Jan;23(1):48-54.
- (284) Nygaard OP, Trumpy JH, Mellgren SI. Recovery of sensory function after surgical decompression in carpal tunnel syndrome. Acta Neurol Scand 1996 Oct;94(4):253-7.
- (285) Oertel J, Schroeder HW, Gaab MR. Dual-portal endoscopic release of the transverse ligament in carpal tunnel syndrome: results of 411 procedures with special reference to technique, efficacy, and complications. Neurosurgery 2006 Aug;59(2):333-40.
- (286) Okutsu I, Hamanaka I, Ninomiya S, Takatori Y, Shimizu K, Ugawa Y. Results of endoscopic management of carpal-tunnel syndrome in long-term haemodialysis versus idiopathic patients. Nephrol Dial Transplant 1993;8(10):1110-4.
- (287) Olney JR, Quenzer DE, Makowsky M. Contested claims in carpal tunnel surgery: outcome study of worker's compensation factors. Iowa Orthop J 1999;19:111-21.
- (288) Ozer H, Solak S, Oguz T, Ocguder A, Colakoglu T, Babacan A. Alkalinisation of local anaesthetics prescribed for pain relief after surgical decompression of carpal tunnel syndrome. J Orthop Surg (Hong Kong) 2005 Dec;13(3):285-9.
- (289) Oztas O, Turan B, Bora I, Karakaya MK. Ultrasound therapy effect in carpal tunnel syndrome. Arch Phys Med Rehabil 1998 Dec;79(12):1540-4.
- (290) Padua L, Padua R, Aprile I, Caliandro P, Tonali P. Boston Carpal Tunnel Questionnaire: the influence of diagnosis on patient-oriented results. Neurol Res 2005 Jul;27(5):522-4.
- (291) Pagnanelli DM, Barrer SJ. Bilateral carpal tunnel release at one operation: report of 228 patients. Neurosurgery 1992 Dec;31(6):1030-3.
- (292) Palmer DH, Paulson JC, Lane-Larsen CL, Peulen VK, Olson JD. Endoscopic carpal tunnel release: a comparison of two techniques with open release. Arthroscopy 1993;9(5):498-508.
- (293) Palmer KT, Harris EC, Coggon D. Carpal tunnel syndrome and its relation to occupation: a systematic literature review. Occup Med (Lond) 2007 Jan;57(1):57-66.
- (294) Park SH, Cho BH, Ryu KS, Cho BM, Oh SM, Park DS. Surgical outcome of endoscopic carpal tunnel release in 100 patients with carpal tunnel syndrome. Minim Invasive Neurosurg 2004 Oct;47(5):261-5.

- (295) Patil S, Ramakrishnan M, Stothard J. Local anaesthesia for carpal tunnel decompression: a comparison of two techniques. J Hand Surg [Br] 2006 Dec;31(6):683-6.
- (296) Patrick DL, Deyo RA. Generic and disease-specific measures in assessing health status and quality of life. Med Care 1989 Mar;27(3 Suppl):S217-S232.
- (297) Pazzaglia C, Caliandro P, Aprile I, Mondelli M, Foschini M, Tonali PA, et al. Multicenter study on carpal tunnel syndrome and pregnancy incidence and natural course. Acta Neurochir Suppl 2005;92:35-9.
- (298) Pierre-Jerome C, Bekkelund SI. Magnetic resonance assessment of the double-crush phenomenon in patients with carpal tunnel syndrome: a bilateral quantitative study. Scand J Plast Reconstr Surg Hand Surg 2003;37(1):46-53.
- (299) Porrata H, Porrata A, Sosner J. New carpal ligament traction device for the treatment of carpal tunnel syndrome unresponsive to conservative therapy. J Hand Ther 2007 Jan;20(1):20-7.
- (300) Posch JL, Prpic I. Surgical treatment of the carpal tunnel syndrome. Handchirurgie 1975;7(2):95-8.
- (301) Pretto Flores L. Endoscopic carpal tunnel release: A comparative study to the conventional open technique. Arq Neuro-Psiquiatr 2005;63(3 A):637-42.
- (302) Raffi GB, Lodi V, Malenchini G, Missere M, Naldi M, Tabanelli S, et al. Cumulative trauma disorders of the upper limbs in workers on an agricultural farm. Arh Hig Rada Toksikol 1996 Mar;47(1):19-23.
- (303) Randolph JA. Carpal tunnel syndrome. Testing the sensitivity and validity of four "localized discomfort" instruments. AAOHN J 2000 Aug;48(8):385-94.
- (304) Removed during review
- (305) Rempel D, Tittiranonda P, Burastero S, Hudes M, So Y. Effect of keyboard keyswitch design on hand pain. J Occup Environ Med 1999 Feb;41(2):111-9.
- (306) Rempel DM, Krause N, Goldberg R, Benner D, Hudes M, Goldner GU. A randomised controlled trial evaluating the effects of two workstation interventions on upper body pain and incident musculoskeletal disorders among computer operators. Occup Environ Med 2006 May;63(5):300-6.
- (307) Roquelaure Y, Raimbeau G, Dano C, Martin YH, Pelier-Cady MC, Mechali S, et al. Occupational risk factors for radial tunnel syndrome in industrial workers. Scand J Work Environ Health 2000 Dec;26(6):507-13.

- (308) Rosales RS, Delgado EB, Diez de la Lastra-Bosch. Evaluation of the Spanish version of the DASH and carpal tunnel syndrome health-related quality-of-life instruments: cross-cultural adaptation process and reliability. J Hand Surg [Am] 2002 Mar;27(2):334-43.
- (309) Sambandam SN, Priyanka P, Gul A, Ilango B. Critical analysis of outcome measures used in the assessment of carpal tunnel syndrome. Int Orthop 2007 Mar 17.
- (310) Sato Y, Honda Y, Iwamoto J, Kanoko T, Satoh K. Amelioration by mecobalamin of subclinical carpal tunnel syndrome involving unaffected limbs in stroke patients. J Neurol Sci 2005 Apr 15;231(1-2):13-8.
- (311) Schafer W, Sander KE, Walter A, Weitbrecht WU. [Agee endoscopic operation of carpal tunnel syndrome in comparison with open surgical technique]. Handchir Mikrochir Plast Chir 1996 May;28(3):143-6.
- (312) Schlenker JD, Koulis CP, Kho LK. Synovialectomy and reconstruction of the retinaculum flexorum in median nerve decompression: technique and early results. Handchir Mikrochir Plast Chir 1993 Mar;25(2):66-71.
- (313) Schonauer F, Belcher HJ. Anthropometry and endoscopic carpal tunnel release. J Hand Surg [Br] 1999 Feb;24(1):6-8.
- (314) Schonauer F, Varma S, Belcher HJ. Endoscopic carpal tunnel release: practice in evolution. Scand J Plast Reconstr Surg Hand Surg 2003;37(6):360-4.
- (315) Schrijver HM, Gerritsen AA, Strijers RL, Uitdehaag BM, Scholten RJ, de Vet HC, et al. Correlating nerve conduction studies and clinical outcome measures on carpal tunnel syndrome: lessons from a randomized controlled trial. J Clin Neurophysiol 2005 Jun;22(3):216-21.
- (316) Sennwald GR, Benedetti R. The value of one-portal endoscopic carpal tunnel release: a prospective randomized study. Knee Surg Sports Traumatol Arthrosc 1995;3(2):113-6.
- (317) Seradge H, Seradge E. Piso-triquetral pain syndrome after carpal tunnel release. J Hand Surg [Am] 1989 Sep;14(5):858-62.
- (318) Serra JM, Benito JR, Monner J. Carpal tunnel release with short incision. Plast Reconstr Surg 1997 Jan;99(1):129-35.
- (319) Sheon RP. Repetitive strain injury. 2. Diagnostic and treatment tips on six common problems. The Goff Group. Postgrad Med 1997 Oct;102(4):72-8, 81, 85.
- (320) Silverstein B, Viikari-Juntura E, Kalat J. Use of a prevention index to identify industries at high risk for work-related musculoskeletal disorders of the neck, back,

- and upper extremity in Washington state, 1990-1998. Am J Ind Med 2002 Mar;41(3):149-69.
- (321) Simpson RL, Fern SA. Multiple compression neuropathies and the double-crush syndrome. Orthop Clin North Am 1996 Apr;27(2):381-8.
- (322) Staal JB, de Bie RA, Hendriks EJ. Aetiology and management of work-related upper extremity disorders. Best Pract Res Clin Rheumatol 2007 Feb;21(1):123-33.
- (323) Stark B, Engkvist-Lofmark C. [Endoscopic operation or conventional open surgical technique in carpal tunnel syndrome: a prospective comparative study]. Handchir Mikrochir Plast Chir 1996 May;28(3):128-32.
- (324) Stasinopoulos D, Stasinopoulos I, Johnson MI. Treatment of carpal tunnel syndrome with polarized polychromatic noncoherent light (Bioptron light): a preliminary, prospective, open clinical trial. Photomed Laser Surg 2005 Apr;23(2):225-8.
- (325) Stransky M, Rubin A, Lava NS, Lazaro RP. Treatment of carpal tunnel syndrome with vitamin B6: a double-blind study. South Med J 1989 Jul;82(7):841-2.
- (326) Szabo RM. Outcomes assessment in hand surgery: when are they meaningful? J Hand Surg [Am] 2001 Nov;26(6):993-1002.
- (327) Tanaka S, Wild DK, Cameron LL, Freund E. Association of occupational and non-occupational risk factors with the prevalence of self-reported carpal tunnel syndrome in a national survey of the working population. Am J Ind Med 1997 Nov;32(5):550-6.
- (328) Thomas RE, Vaidya SC, Herrick RT, Congleton JJ. The effects of biofeedback on carpal tunnel syndrome. Ergonomics 1993 Apr;36(4):353-61.
- (329) Thomsen JF, Hansson GA, Mikkelsen S, Lauritzen M. Carpal tunnel syndrome in repetitive work: a follow-up study. Am J Ind Med 2002 Oct;42(4):344-53.
- (330) Tittiranonda P, Rempel D, Armstrong T, Burastero S. Effect of four computer keyboards in computer users with upper extremity musculoskeletal disorders. Am J Ind Med 1999 Jun;35(6):647-61.
- (331) Trumble TE, Gilbert M, McCallister WV. Endoscopic versus open surgical treatment of carpal tunnel syndrome. Neurosurg Clin N Am 2001 Apr;12(2):255-66.
- (332) Tsai CP, Liu CY, Lin KP, Wang KC. Efficacy of botulinum toxin type a in the relief of Carpal tunnel syndrome: A preliminary experience. Clin Drug Investig 2006;26(9):511-5.

- (333) Turner JA, Franklin G, Fulton-Kehoe D, Egan K, Wickizer TM, Lymp JF, et al. Prediction of chronic disability in work-related musculoskeletal disorders: a prospective, population-based study. BMC Musculoskelet Disord 2004 May 24;5:14.
- (334) Tuzuner S, Sherman GM, Ozkaynak S, Ozcanli H. Endoscopic carpal tunnel release: modification of Menon's technique and data from 191 cases. Arthroscopy 2004 Sep;20(7):721-7.
- (335) van den Bekerom MP, Breemans E, Schaffer K. Outcome of open versus endoscopic approach for the surgical treatment of carpal tunnel syndrome. Acta Orthop Belg 2006 Jun;72(3):288-95.
- (336) Vasen AP, Kuntz KM, Simmons BP, Katz JN. Open versus endoscopic carpal tunnel release: a decision analysis. J Hand Surg [Am] 1999 Sep;24(5):1109-17.
- (337) Verhagen AP, Karels C, Bierma-Zeinstra SM, Burdorf L, Feleus A, Dahaghin S, et al. Ergonomic and physiotherapeutic interventions for treating work-related complaints of the arm, neck or shoulder in adults. Cochrane Database Syst Rev 2006;3:CD003471.
- (338) Vossinakis IC, Stavroulaki P, Paleochorlidis I, Badras LS. Reducing the pain associated with local anaesthetic infiltration for open carpal tunnel decompression. J Hand Surg [Br] 2004 Aug;29(4):399-401.
- (339) Wang AA, Whitaker E, Hutchinson DT, Coleman DA. Pain levels after injection of corticosteroid to hand and elbow. Am J Orthop 2003 Aug;32(8):383-5.
- (340) Ware JE, Jr., Sherbourne CD. The MOS 36-item short-form health survey (SF-36). I. Conceptual framework and item selection. Med Care 1992 Jun;30(6):473-83.
- (341) Weber RA, Rude MJ. Clinical outcomes of carpal tunnel release in patients 65 and older. J Hand Surg [Am] 2005 Jan;30(1):75-80.
- (342) Weintraub MI, Cole SP. Pulsed magnetic field therapy in refractory carpal tunnel syndrome: Electrodiagnostic parameters Pilot study. J Back Musculoskelet Rehabil 2005;18(3-4):79-83.
- (343) Weintraub MI, Cole SP. Time-Varying, Biaxial Magnetic Stimulation in Refractory Carpal Tunnel Syndrome: A Novel Approach. A Pilot Study. Semin Integr Med 2005;3(4):123-8.
- (344) Wellman H, Davis L, Punnett L, Dewey R. Work-related carpal tunnel syndrome (WR-CTS) in Massachusetts, 1992-1997: source of WR-CTS, outcomes, and employer intervention practices. Am J Ind Med 2004 Feb;45(2):139-52.

- (345) Werner RA, Franzblau A, Gell N. Randomized controlled trial of nocturnal splinting for active workers with symptoms of carpal tunnel syndrome. Arch Phys Med Rehabil 2005 Jan;86(1):1-7.
- (346) Werner RA. Evaluation of work-related carpal tunnel syndrome. J Occup Rehabil 2006 Jun;16(2):207-22.
- (347) Wewers ME, Lowe NK. A critical review of visual analogue scales in the measurement of clinical phenomena. Res Nurs Health 1990 Aug;13(4):227-36.
- (348) Wilbourn AJ, Gilliatt RW. Double-crush syndrome: a critical analysis. Neurology 1997 Jul;49(1):21-9.
- (349) Wong SM, Hui AC, Tang A, Ho PC, Hung LK, Wong KS, et al. Local vs systemic corticosteroids in the treatment of carpal tunnel syndrome. Neurology 2001 Jun 12;56(11):1565-7.
- (350) Wood VE, Biondi J. Double-crush nerve compression in thoracic-outlet syndrome. J Bone Joint Surg Am 1990 Jan;72(1):85-7.
- (351) Yagci I, Ucan H, Yilmaz L, Yagmurlu F, Keskin D, Bodur H. Comparison of splinting, splinting plus local steroid injection and surgery in carpal tunnel syndrome treatment. Turk Fiz Tip Rehab Derg 2006;52(2):55-60.
- (352) Yoshida A, Okutsu I, Hamanaka I, Motomura T. Results of endoscopic management of primary versus recurrent carpal tunnel syndrome in long-term haemodialysis patients. Hand Surg 2004 Dec;9(2):165-70.
- (353) 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 Dec;26(6):657-60.
- (354) Zyluk A, Strychar J. A comparison of two limited open techniques for carpal tunnel release. J Hand Surg [Br] 2006 Oct;31(5):466-72.

IV.APPENDIXES

APPENDIX I: LITERATURE SEARCHES

DATABASES SEARCHED

The search for eligible literature began with a search for applicable systematic reviews. The search for systematic reviews was performed using the following databases. The full search strategies are displayed below.

- MEDLINE (from 1966 through April 6, 2007)
- EMBASE (from 1966 through April 6, 2007)
- The Cochrane Database of Systematic Reviews (through April 6, 2007)

This initial search yielded 109 systematic reviews, of which 51 were retrieved and evaluated. Fifty-eight systematic reviews were not retrieved because their titles indicated they reviewed topics that were irrelevant to the recommendations in this guideline. Of the fifty-one retrieved, five systematic reviews met all inclusion criteria. These systematic reviews were updated with controlled trials identified through MEDLINE and EMBASE searches.

The literature searches for recommendations that were not addressed by existing systematic reviews were performed using one or more of the same databases identified previously except through June 12, 2007. A search of the CINAHL database from 1982 through June 12, 2007 was also conducted for Recommendation 9.

All literature searches were supplemented with manual screening of bibliographies in publications accepted for inclusion into the evidence base. In addition, the bibliographies of recent review articles were searched for potentially relevant citations.

SEARCH STRATEGIES

ORIGINAL SEARCH FOR SYSTEMATIC REVIEWS

Our search for systematic reviews using PubMed included the following search strategy, with limits of publication dates 1966 to present, English language, and humans:

(("carpal tunnel syndrome"[TIAB] NOT Medline[SB]) OR "carpal tunnel syndrome"[MeSH Terms] OR carpal tunnel[Text Word]) AND systematic[sb]

Our search for systematic reviews using EMBASE included the following search strategy with limits of publication dates 1966 to present, English language, and humans:

carpal AND tunnel AND ([cochrane review]/lim OR [systematic review]/lim) AND [humans]/lim AND [embase]/lim)

SEARCHES FOR PRIMARY STUDIES FROM SYSTEMATIC REVIEWS

The following search strategies are the searches we used to update the identified systematic review. In all cases, we replicated as closely as possible the strategies identified by the original authors of the applicable systematic review. These strategies however may not be precisely duplicated due to lack of complete information in the original systematic review.

MARSHALL ET AL

- 1. clinical trial.pt.
- 2. randomized controlled trial.pt.
- 3. tu.fs.
- 4. dt.fs.
- 5. random\\$.tw.
- 6. (double and blind\$).tw.
- 7. placebo\$.tw.
- 8. exp Comparative Study/
- 9. 1 or 2 or 3 or 4 or 5 or 6 or 7 or 8
- 10. exp Carpal tunnel syndrome/
- 11. exp Steroids/
- 12. exp injections/ or exp injections, intra-articular/
- 13. 11 or 12
- 14. 10 and 13
- 15. 9 and 14

O'CONNOR ET AL

- 1. randomized controlled trial.pt.
- 2. randomized controlled trials/
- 3. controlled clinical trial.pt.
- 4. controlled clinical trials/
- 5. random allocation/
- 6. double-blind method/
- 7. single-blind method/

- 8. clinical trial.pt.
- 9. exp clinical trials/
- 10. (clin\$ adj25 trial\$).tw.
- 11. ((singl\$ or doubl\$ or tripl\$ or trebl\$) adj25 (blind\$ or mask\$ or dummy)).tw.
- 12. placebos/
- 13. placebo\$.tw.
- 14. random\$.tw.
- 15. research design/
- 16. (clinical trial phase i or clinical trial phase ii or clinical trial phase iii or clinical trial phase iv).pt.
- 17. multicenter study.pt.
- 18. meta analysis.pt.
- 19. prospective studies/
- 20. intervention studies/
- 21. cross-over studies/
- 22. meta-analysis/
- 23. (meta?analys\$ or systematic review\$).tw.
- 24. control\$.tw.
- 25 or/1-24
- 26. human/
- 27. 25 and 26
- 28. Carpal tunnel syndrome/dt,rh,th [Drug Therapy, Rehabilitation, Therapy]
- 29. 27 and 28

SCHOLTEN ET AL

Specific search for CTS: 'carpal tunnel syndrome [mesh]' OR 'carpal tunnel syndrome [tw]' OR 'carpal tunnel [tw]' OR 'carp* syndr* [tw]' OR 'carp* tunn* [tw]' OR 'tunn* syndr* [tw]' OR 'median nerve entrapment [mesh]' OR 'median nerve entrapment [tw]'. Specific search for surgical interventions: 'surgical [mesh]' OR 'surgical [tw]' OR 'surgery [mesh]' OR 'surgery [tw]' OR 'release [tw]' OR 'reconstruct* [tw]' OR 'epineurotomy [tw]'.

VERDUGO ET AL

- 1. ('median nerve entrapment' or 'carpal tunnel syndrome' or 'entrapment neuropathy')
- 2. limit 1 to english language
- 3. limit 2 to yr="2003 2007"
- 4. and randomized.mp.
- 5. remove duplicates from 4
- 6. from 5 keep 98-127

SEARCH STRATEGIES FOR RECOMMENDATIONS WITHOUT RELEVANT SYSTEMATIC REVIEWS

RECOMMENDATION 3

In addition to the associated conditions found in the systematic reviews and relevant RCT's the Work Group examined coexisting cervical radiculopathy and CTS in the workplace. Searches for applicable studies concerning coexisting cervical radiculopathy and carpal tunnel syndrome were limited to what has been termed "double crush syndrome" (in conducting this search we acknowledge that there is controversy surrounding the existence of "double crush syndrome"). The searches for coexisting cervical radiculopathy and CTS in the workplace were conducted as follows:

PubMed and Embase databases were searched for the phrase "double crush" [All Fields].

The PubMed database was searched using the following strategy: (carpal tunnel or carpal tunnel syndrome or median neuropathy) AND work AND (intervention OR activity OR activities OR moderation OR behavior OR modification OR modify OR restriction*) AND English[lang]

RECOMMENDATION 7

After examining the included studies from controlled trials examining surgical or postsurgical treatments, another literature search was conducted. All study designs focusing on surgical and post-surgical treatments were examined for reports of infection.

PubMed:

"Carpal Tunnel Syndrome/surgery" [Majr] OR "Median Neuropathy/surgery" [Majr] NOT Comment [Publication Type] NOT Letter [Publication Type] NOT Biography [Publication Type] NOT Historical Article [Publication Type] NOT Practice Guideline [Publication Type] NOT Guideline [Publication Type] NOT Case Reports [Publication Type] NOT Editorial [Publication Type] NOT Clinical Trial [Publication Type] NOT Meta-Analysis [Publication Type] NOT Review [Publication Type] NOT Validation Studies [Publication Type] AND (hasabstract [text] AND (Humans [Mesh]) AND (English [lang]))

RECOMMENDATION 8

The PubMed database was searched for carpal tunnel syndrome and postoperative or rehabilitation studies. The search was limited to clinical trials only using the following strategy:

((("carpal tunnel syndrome"[TIAB] NOT Medline[SB]) OR "carpal tunnel syndrome"[MeSH Terms] OR carpal tunnel[Text Word]) OR ("carpal tunnel syndrome"[MeSH Terms] OR carpal tunnel syndrome[Text Word]) OR ("median neuropathy"[MeSH Terms] OR median neuropathy[Text Word])) AND ((("postoperative period"[TIAB] NOT Medline[SB]) OR "postoperative period"[MeSH Terms] OR postoperative[Text Word] OR "postoperative care"[MeSH Terms] OR postoperative care[Text Word]) OR ("rehabilitation"[Subheading] OR "rehabilitation"[MeSH Terms]

OR rehabilitation[Text Word])) AND (English[lang] AND (Clinical Trial[ptyp] OR Clinical Trial, Phase II[ptyp] OR Clinical Trial, Phase III[ptyp] OR Clinical Trial, Phase IV[ptyp]))

RECOMMENDATION 9

The PubMed database was searched using the following strategy: (("michigan hand outcome" OR "michigan hand outcomes" OR "Boston Carpal Tunnel Questionnaire" OR "Boston Questionnaire" OR "global symptom score" OR "Multidimensional Health Assessment Questionnaire" OR "visual analogue scale" OR "visual analog scale" OR Disability of the Arm, Shoulder, and Hand OR "Carpal Tunnel Syndrome Instrument" OR "symptom severity score" OR "functional status score" OR "short form 36" OR "SF36" OR "SF-12" OR "SF12" OR "Levine functional score" OR "Brigham and Women's carpal tunnel questionnaire" OR "brief pain inventory" OR postoperative pain OR return to work OR work absence) AND (median neuropathy OR carpal tunnel OR carpal tunnel syndrome)) AND English[lang]

The EMBASE database was searched using the following strategy: ('michigan hand outcome' OR 'michigan hand outcomes' OR 'boston carpal tunnel questionnaire' OR 'boston questionnaire' OR 'global symptom score' OR 'multidimensional health assessment questionnaire' OR 'visual analogue scale' OR 'visual analog scale' OR 'disability of the arm, shoulder, and hand' OR 'carpal tunnel syndrome instrument' OR 'symptom severity score' OR 'functional status score' OR 'short form 36' OR 'sf36' OR 'sf-12' OR 'sf12' OR 'levine functional score' OR 'brigham and womens carpal tunnel questionnaire' OR 'brief pain inventory' OR postoperative AND pain OR return AND to AND work OR work AND absence) AND ('carpal tunnel syndrome'/exp OR 'median neuropathy'/exp) AND [embase]/lim

The CINAHL database was searched using the following strategy: (michigan hand outcome OR michigan hand outcomes OR boston carpal tunnel questionnaire OR boston questionnaire OR global symptom score OR multidimensional health assessment questionnaire OR visual analogue scale OR visual analog scale OR disability of the arm, shoulder, and hand OR carpal tunnel syndrome instrument OR symptom severity score OR functional status score OR short form 36 OR sf36 OR sf-12 OR sf12 OR levine functional score OR brigham and womens carpal tunnel questionnaire OR brief pain inventory OR postoperative pain OR return to work OR work absence) AND (carpal tunnel syndrome or median neuropathy)

APPENDIX II: ARTICLE INCLUSIONS AND EXCLUSIONS

Flow charts illustrating study attrition are depicted in Figure 1 - Figure 10 below. All abstracts were downloaded, reviewed, and evaluated for the following criteria:

EXCLUSION CRITERIA

- Abstracts and unpublished study reports.
- Cadaveric, animal or in vitro studies.
- Letters, case reports, historical articles, editorials, and commentaries.
- Non prospective studies.
- Studies where gender is restricted.
- Studies where results for CTS population cannot be separated from results from other populations.
- Studies with < 10 patients.
- Studies with patients under 18 years of age.
- Studies written in languages other than English.

INCLUSION CRITERIA

- Studies evaluating a treatment or intervention for CTS.
- Studies that measured the validity, reliability, or responsiveness of any assessment instrument.
- The following study designs: randomized controlled trials or prospective controlled trials. Where appropriate, observational study designs were also considered (i.e. prospective cohorts, case series, etc.).
- Studies where data can be extracted for statistical analysis.
- Studies reporting patient-oriented outcome measures using previously validated instruments.
- Studies that diagnose CTS with electro-diagnostic tests, signs and/or symptoms of the syndrome.

Full articles were retrieved for all abstracts meeting these criteria. Once retrieved the complete articles were reviewed and evaluated for inclusion. See Figure 1 - Figure 10 below

INCLUDED AND EXCLUDED ARTICLES FLOWCHART

RECOMMENDATIONS 1,2,3,4

Figure 1

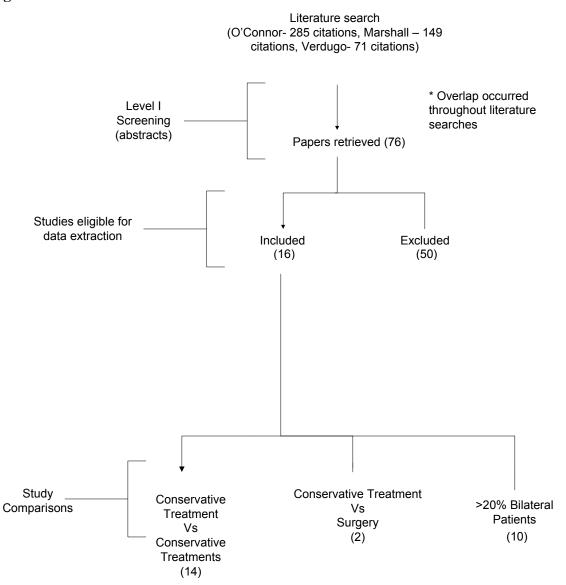


Figure 2
O'Connor Systematic Review Update

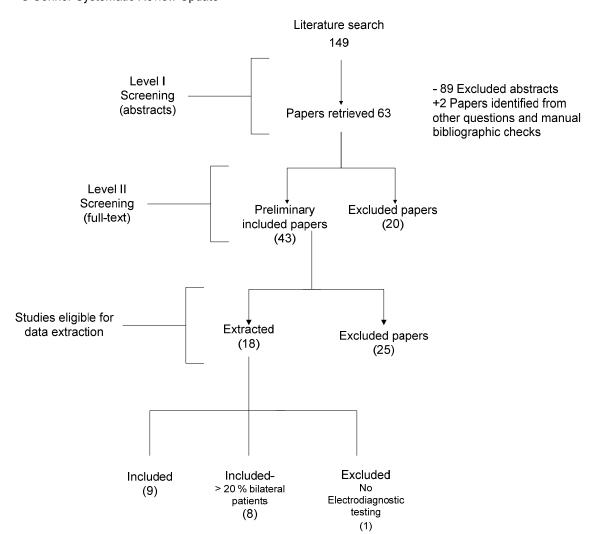


Figure 3

Marshall Systematic Review Update

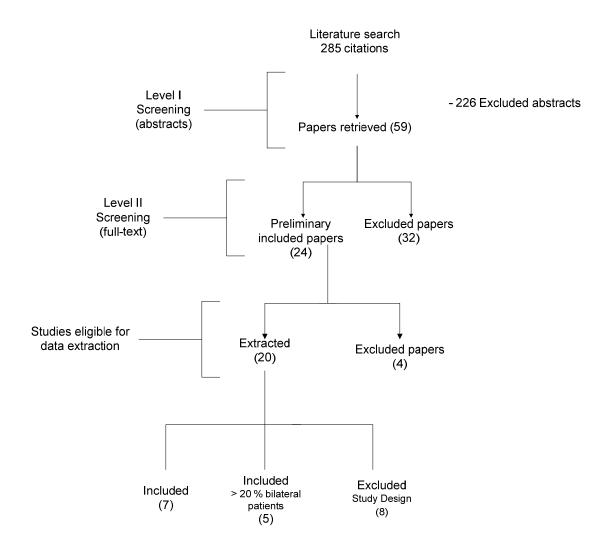
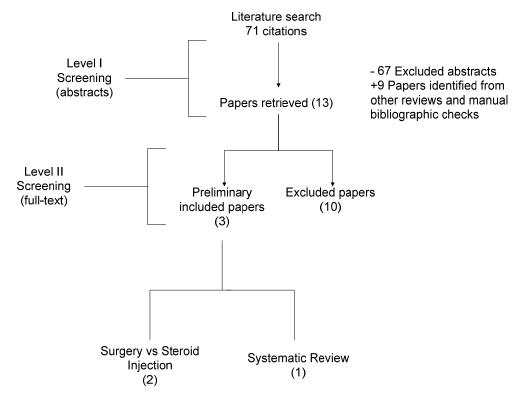


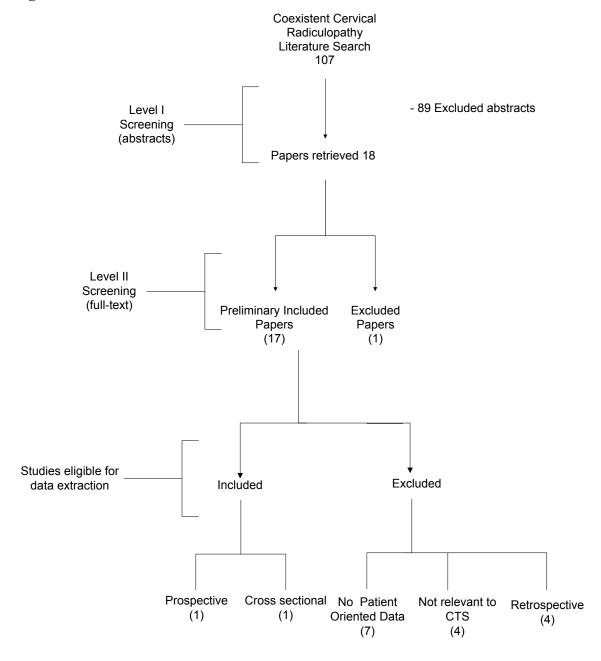
Figure 4

Update to Verdugo Systematic Review



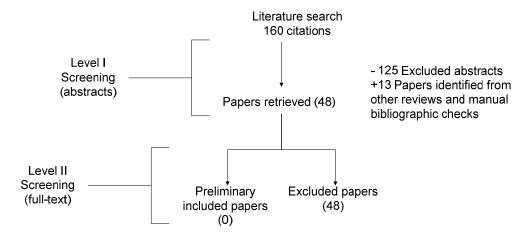
COEXISTENT CERVICAL RADICULOPATHY

Figure 5



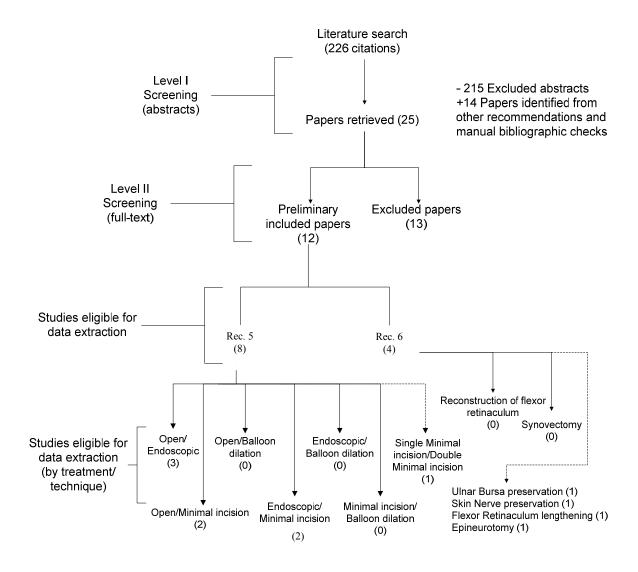
CTS IN THE WORKPLACE

Figure 6



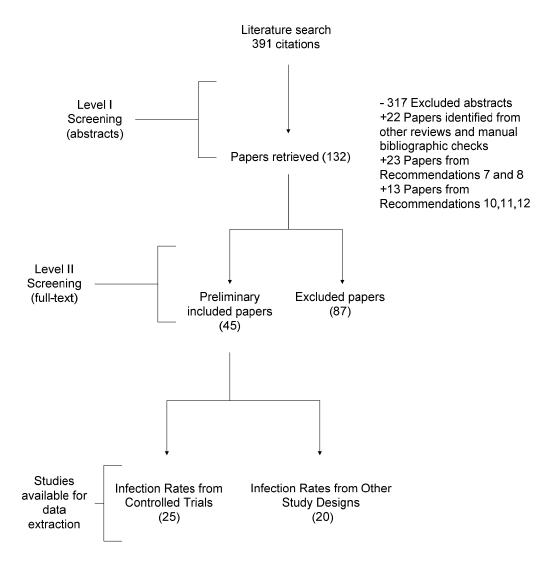
RECOMMENDATION 5 AND 6

Figure 7



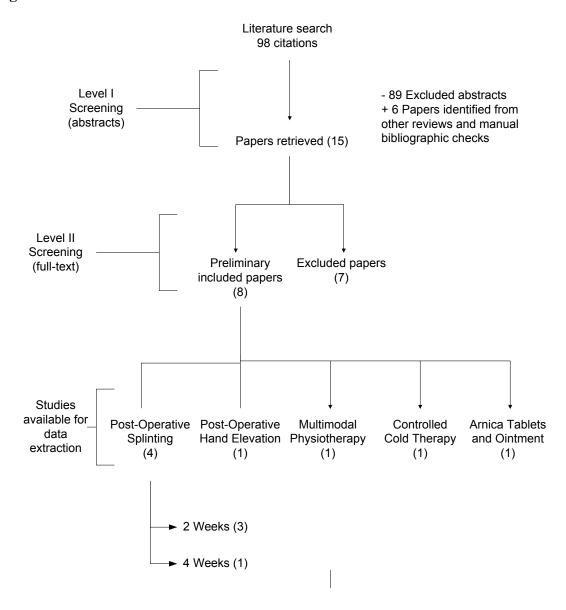
RECOMMENDATION 7

Figure 8



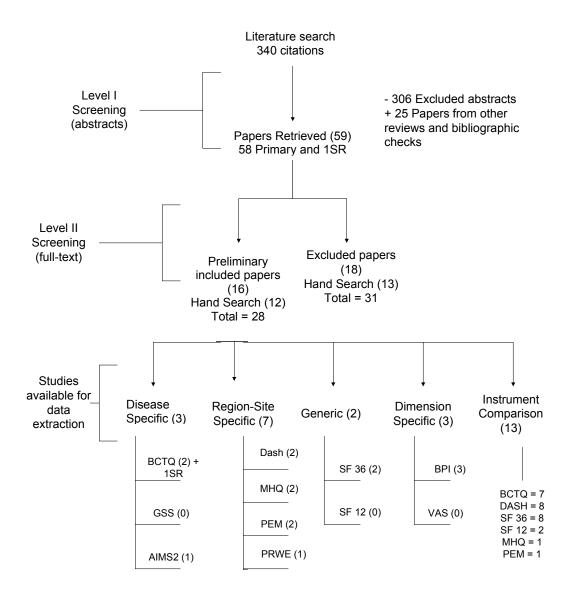
RECOMMENDATION 8

Figure 9



RECOMMENDATION 9

Figure 10



APPENDIX III: RATING EVIDENCE QUALITY

We considered the quality of the available evidence when grading the strength of guideline recommendations. Quality was determined using a "Levels of Evidence" approach in which five levels of evidence were designed for each of four study designs; therapeutic, prognostic, diagnostic and economic or decision modeling. The higher the level of evidence, the greater the ability to draw causal inferences from the results of a study and, hence, the greater the quality of that study.

Levels of Evidence For Primary Research Question¹

September 28, 2004

	Types of Studies				
	Therapeutic Studies – Investigating the results of treatment	Prognostic Studies – Investigating the effect of a patient characteristic on the outcome of disease	Diagnostic Studies – Investigating a diagnostic test	Economic and Decision Analyses – Developing an economic or decision model	
Level I	High quality randomized trial with statistically significant difference or no statistically significant difference but narrow confidence intervals Systematic Review ² of Level I RCTs (and study results were homogenous ³)	High quality prospective study ⁴ (all patients were enrolled at the same point in their disease with ≥ 80% follow-up of enrolled patients) Systematic review ² of Level I studies	Testing of previously developed diagnostic criteria on consecutive patients (with universally applied reference "gold" standard) Systematic review of Level I studies	Sensible costs and alternatives; values obtained from many studies; with multiway sensitivity analyses Systematic review ² of Level I studies	
Level II	Lesser quality RCT (e.g. < 80% follow- up, no blinding, or improper randomization) Prospective ⁴ comparative study ⁵ Systematic review ² of Level II studies or Level 1 studies with inconsistent results	Retrospective ⁶ study Untreated controls from an RCT Lesser quality prospective study (e.g. patients enrolled at different points in their disease or <80% follow-up.) Systematic review ² of Level II studies	Development of diagnostic criteria on consecutive patients (with universally applied reference "gold" standard) Systematic review ² of Level II studies	Sensible costs and alternatives; values obtained from limited studies; with multiway sensitivity analyses Systematic review ² of Level II studies	
Level III	Case control study ⁷ Retrospective ⁶ comparative study ⁵ Systematic review ² of Level III studies	• Case control study ⁷	Study of non-consecutive patients; without consistently applied reference "gold" standard Systematic review ² of Level III studies	Analyses based on limited alternatives and costs; and poor estimates Systematic review ² of Level III studies	
Level IV	Case Series ⁸	Case series	Case-control study Poor reference standard	Analyses with no sensitivity analyses	
Level V	Expert Opinion	Expert Opinion	Expert Opinion	Expert Opinion	

- 1. A complete assessment of quality of individual studies requires critical appraisal of all aspects of the study design.
- A combination of results from two or more prior studies.
- Studies provided consistent results.
- 4. Study was started before the first patient enrolled.
- 5. Patients treated one way (e.g. cemented hip arthroplasty) compared with a group of patients treated in another way (e.g. uncemented hip arthroplasty) at the same institution.
- 6. The study was started after the first patient enrolled.
- Patients identified for the study based on their outcome, called "cases"; e.g. failed total arthroplasty, are compared to those who did
- not have outcome, called "controls"; e.g. successful total hip arthroplasty.

 Patients treated one way with no comparison group of patients treated in another way.

APPENDIX IV: EVIDENCE TABLES

SEE EVIDENCE TABLES DOCUMENT (EVIDENCE TABLES.PDF)

Please refer to the accompanying PDF Document ...

LIST OF TABLES

Table 1. Systematic Review (and Meta Analysis) Bibliographic Information	I
Table 2. Non-Operative/Conservative Treatment Excluded Articles	5
Table 3. Non-Operative/Conservative Treatment Relevant Articles from Systematic Reviews	9
Table 4. Non-Operative/Conservative Treatment Included Articles	10
Table 5. Non-Operative/Conservative Treatment Study Design and Quality	11
Table 6. Non-Operative/Conservative Treatment Patient Characteristics	12
Table 7. Non-Operative/Conservative Treatment Study Inclusion/Exclusion Criteria	16
Table 8. Non-Operative/Conservative Treatment Workers Compensation	19
Table 9. Non-Operative/Conservative Treatment Continuous Data Results	22
Table 10. Non-Operative/Conservative Treatment Dichotomous Data Results	43
Table 11. Non-Operative/Conservative Treatment (bilateral) Relevant Articles from Systematic Reviews	45
Table 12. Non-Operative/Conservative Treatment (bilateral) Included Articles	46
Table 13. Non-Operative/Conservative Treatment (bilateral) Study Design and Quality	47
Table 14. Non-Operative/Conservative Treatment (bilateral) Patient Characteristics	48
Table 15. Non-Operative/Conservative Treatment (bilateral) Study Inclusion/Exclusion Criteria	
Table 16. Non-Operative/Conservative Treatment (bilateral) Workers Compensation	
Table 17. Non-Operative/Conservative Treatment (bilateral) Continuous Data Results	
Table 18. Non-Operative/Conservative Treatment (bilateral) Dichotomous Data Results	
Table 19. Comparisons for Non-Operative/Conservative Treatment	
Table 20. Benefits and Harms for Non-Operative/Conservative Treatment	
Table 21. Minimal Detectable Effect Size for Non-Operative/Conservative Treatment	
Table 22. Work-Related CTS Excluded Articles	
Table 23. Surgical Treatment Excluded Articles	
Table 24. Surgical Treatment Relevant Articles from Systematic Reviews	
Table 25. Surgical Treatment Included Articles	
Table 26. Surgical Treatment Study Design and Quality	
Table 27. Surgical Treatment Patient Characteristics	
Table 28. Surgical Treatment Study Inclusion/Exclusion Criteria	
Table 29. Surgical Treatment Workers Compensation	
Table 30. Surgical Treatment Continuous Data Results	
Table 31. Surgical Treatment Dichotomous Data Results	
Table 32. Benefits and Harms for Surgical Treatment - Minimal Incision vs. OCTR	
Table 33. Minimal Detectable Effect Size for Surgical Treatment - Minimal Incision vs. OCTR	
Table 34. Benefits and Harms for Surgical Treatment - Minimal Incision vs. ECTR	
Table 35. Minimal Detectable Effect Size for Surgical Treatment - Minimal Incision vs. ECTR	
Table 36. Benefits and Harms for Surgical Treatment - Adjuvant Methods	
Table 37. Minimal Detectable Effect Size for Surgical Treatment - Adjuvant Methods	
Table 38. Pre-operative Antibiotics Excluded Articles	
Table 39 Pre-operative Antibiotics Included Articles	144

Table 40. Pre-operative Antibiotics Infection Rates from Controlled Trials	
Table 41. Pre-operative Antibiotics Infection Rates from Other Study Designs	148
Table 42. Post-operative Treatment Excluded Articles	149
Table 43. Post-operative Treatment Included Articles	150
Table 44. Post-operative Treatment Study Design and Quality	151
Table 45. Post-operative Treatment Patient Characteristics	152
Table 46. Post-operative Treatment Study Inclusion/Exclusion Criteria	154
Table 47. Post-operative Treatment Workers Compensation	156
Table 48. Post-operative Treatment Continuous Data Results	157
Table 49. Post-operative Treatment Dichotomous Data Results	160
Table 50. Benefits and Harms for Post-operative Treatment	162
Table 51. Minimal Detectable Effect Size for Post-operative Treatment	163
Table 52. Instrument Evaluation Excluded Articles	166
Table 53. Instrument Evaluation Included Articles	168
Table 54. Instrument Evaluation Summary of Instruments	170
Table 55. Instrument Evaluation Study Design and Patient Characteristics	172
Table 56. Instrument Evaluation Quality	173
Table 57. Instrument Evaluation Funding	175
Table 58. Instrument Evaluation BCTQ Summary	178
Table 59. Instrument Evaluation AIMS2 Summary	179
Table 60. Instrument Evaluation DASH Summary	180
Table 61. Instrument Evaluation MHQ Summary	181
Table 62. Instrument Evaluation PEM Summary	182
Table 63. Instrument Evaluation VAS Summary	183
Table 64. Instrument Evaluation SF-36 Summary	184
Table 65. Instrument Evaluation SF-12 Summary	185
Table 66. Instrument Evaluation BCTQ Reliability	186
Table 67. Instrument Evaluation DASH Reliability	187
Table 68. Instrument Evaluation PEM Reliability	188
Table 69. Instrument Evaluation BCTQ Convergent Validity	189
Table 70. Instrument Evaluation BCTQ Discriminant Validity	191
Table 71. Instrument Evaluation BCTQ Summary Construct Validity	192
Table 72. Instrument Evaluation BCTQ-SSS Summary Construct Validity	193
Table 73. Instrument Evaluation BCTQ-FSS Summary Construct Validity	194
Table 74. Instrument Evaluation AIMS2 Convergent Validity	195
Table 75. Instrument Evaluation DASH Convergent Validity	
Table 76. Instrument Evaluation DASH Discriminant Validity	199
Table 77. Instrument Evaluation DASH Summary Construct Validity	200
Table 78. Instrument Evaluation PEM Convergent Validity	201
Table 79. Instrument Evaluation PEM Discriminant Validity	202
Table 80. Instrument Evaluation PEM Summary Construct Validity	203
Table 81. Instrument Evaluation VAS Convergent Validity	204
Table 82. Instrument Evaluation VAS Discriminant Validity	205
Table 83. Instrument Evaluation VAS Summary Construct Validity	206
Table 84. Instrument Evaluation SF-36 Convergent Validity	
Table 85 Instrument Evaluation SF-36 Discriminant Validity	211

Table 86. Instrument Evaluation BCTQ Responsiveness	214
Table 87. Instrument Evaluation BCTQ Responsiveness - Variance	215
Table 88. Instrument Evaluation AIMS2 Responsiveness	217
Table 89. Instrument Evaluation AIMS2 Responsiveness - Variance	218
Table 90. Instrument Evaluation DASH Responsiveness	219
Table 91. Instrument Evaluation DASH Responsiveness - Variance	220
Table 92. Instrument Evaluation MHQ Responsiveness	221
Table 93. Instrument Evaluation MHQ Responsiveness - Variance	222
Table 94. Instrument Evaluation PEM Responsiveness	223
Table 95. Instrument Evaluation PEM Responsiveness - Variance	224
Table 96. Instrument Evaluation VAS Responsiveness	225
Table 97. Instrument Evaluation VAS Responsiveness - Variance	226
Table 98. Instrument Evaluation SF-36 Responsiveness	227
Table 99. Instrument Evaluation SF-36 Responsiveness - Variance	229
Table 100. Instrument Evaluation SF-12 Responsiveness	232
Table 101. Instrument Evaluation SF-12 Responsiveness - Variance	

APPENDIX V: CONFLICTS OF INTEREST

All members of the physician Work Group disclosed any conflicts of interest prior to the development of the recommendations for this guideline. Conflicts of interest are disclosed in writing with the American Academy of Orthopaedic Surgeons via a private on-line reporting database and also verbally at the recommendation approval meeting. No member of the CTS Work Group disclosed a conflict of interest for this guideline.

APPENDIX VI: DOCUMENTATION OF APPROVAL

AAOS Work Group Draft Completed December 2007

Outside Specialty Review Panel Comments Completed April 25, 2008

Public Commentary Completed May 2008

AAOS Guidelines and Technology Oversight Committee June 11, 2008

AAOS Evidence Based Practice Committee June 19, 2008

AAOS Council on Research Quality Assessment July 9, 2008

and Technology

AAOS Board of Directors September 12, 2008

A total of 198 professionals were provided the opportunity to comment on the contents of this document during the peer review, public commentary and approval process.

Suggested Citation for referencing this document:

American Academy of Orthopaedic Surgeons Clinical Practice Guideline on Treatment of Carpal Tunnel Syndrome. Rosemont (IL): American Academy of Orthopaedic Surgeons (AAOS); 2008

APPENDIX VII: ADVISORY REVIEW PANEL

Participation in the AAOS peer review process does not constitute an endorsement of this guideline by the participating organization.

Peer review of the draft guideline is completed by an outside Peer Review Advisory Panel. Outside Advisory Panels are convened for each AAOS guideline and consist of experts in the guideline's topic area. These experts represent professional societies other than AAOS and are nominated by the guideline Work Group prior to beginning work on the guideline. For this guideline, thirteen outside peer review organizations were invited to review the draft guideline and all supporting documentation. Eight societies participated in the review of the CTS Treatment guideline draft and seven consented to be listed as a peer review organization in this appendix. One organization requested that the organization name be withheld from publication. The organizations that reviewed the document and consented to publication are listed below:

The American Academy of Neurology (AAN)

The American Academy of Physical Medicine and Rehabilitation (AAPMR)

The American Association of Neurological Surgeons/Congress of Neurological Surgeons (AANS/CNS)

The American Association of Neuromuscular and Electrodiagnostic Medicine (AANEM)

The American College of Occupational and Environmental Medicine (ACOEM)

The American Medical Association (AMA)

The American Society of Plastic Surgeons (ASPS)

Again, participation in the AAOS guideline peer review process does not constitute an endorsement of the guideline by the participating organizations listed above.