



# Effectiveness of TECAR Therapy on Clinical Symptoms and Neurophysiological Parameters of Patients with Carpal Tunnel Syndrome: A Randomized Clinical Trial

Babak Vahdatpour<sup>1</sup>, Hamid Reza Ghasemi<sup>1</sup> and Parisa Taheri<sup>1,\*</sup>

<sup>1</sup>Department of Physical Medicine and Rehabilitation, School of Medicine, Isfahan University of Medical Sciences, Isfahan, Iran

\*Corresponding author: Department of Physical Medicine and Rehabilitation, School of Medicine, Isfahan University of Medical Sciences, Isfahan, Iran. Email: prs\_taheri@yahoo.com

Received 2022 December 17; Revised 2023 February 14; Accepted 2023 February 28.

## Abstract

**Background:** Carpal tunnel syndrome (CTS) is the most typical peripheral entrapment neuropathy. To date, various non-invasive methods have been used to manage CTS. This study investigates the effectiveness of Transfer Energy Capacitive and Resistive (TECAR), a new physical agent modality, on symptoms and neurophysiological parameters in patients with CTS.

**Methods:** In this single-blind randomized clinical trial, 29 patients with mild or moderate CTS were randomly assigned to two groups. Group 1 received TECAR therapy with a frequency of 500 Hz and an intensity of 30 to 50%, 2 sessions/week, in addition to wrist splints and vitamin B supplements for 4 weeks. Group 2 also received a wrist splint and vitamin B for 4 weeks without TECAR therapy. The Visual Analog Scale (VAS), Boston Carpal Tunnel Questionnaire (BCTQ-SSS and FSS), as well as clinical and neurophysiological findings, were assessed before and 8 weeks after treatment. Within and between-group comparisons were made after the intervention period.

**Results:** VAS and the Boston Carpal Tunnel Questionnaire (FSS and SSS) showed significant changes. Clinical CTS tests and electrophysiological parameters did not show significant differences before and after the study. From a clinical perspective (pain relief and functional improvement), the TECAR therapy group showed more definite changes.

**Conclusions:** TECAR can be considered an effective non-invasive treatment for patients with mild to moderate CTS.

**Keywords:** Carpal Tunnel Syndrome, Physical Therapy Modalities, Nerve Conduction Study

## 1. Background

Carpal tunnel syndrome is a common cause of upper extremity disability (1). It is a peripheral entrapment neuropathy because of the involvement of the median nerve in the wrist region.

The symptoms include pain, tingling, paresthesia, and discomfort, and as the disease progresses, hand movement will become impaired (2), paresthesia becomes more severe while driving, and symptoms are relieved by shaking the hand (3).

With a prevalence of 4% in the population, carpal tunnel syndrome is one of the most common entrapment neuropathy (4). The causes of CTS include medical conditions (thyroid disease, kidney failure, acromegaly, etc.), inflammatory disease (lupus, rheumatoid arthritis, etc.), tumors and pseudo-tumors (ganglions, etc.), anatomical anomalies, and in most cases, it is idiopathic (5).

The disease occurs with increased pressure in the carpal tunnel, especially at the level of the carpal ligament,

which results in pressure on the median nerve, decreased epineural blood supply to the nerve, and progression of nerve damage due to local ischemia (6).

Neurophysiologic studies (such as nerve conduction studies) are highly sensitive diagnostic tests that investigate median nerve involvement (7). The degree of a patient's symptoms and level of disability can be assessed using reliable patient-centered tools like the Boston Carpal Tunnel Syndrome Questionnaire (BCTQ). The correlation between clinical findings and nerve conduction study findings allows for assessing the patient's level of disability (8).

Both surgical and non-surgical alternatives are available for treating carpal tunnel syndrome. Physical agent modalities are one of the various non-surgical treatments for carpal tunnel syndrome (7, 9).

Diathermy is one of these modes. Physical therapy uses diathermy, a deep heat current source that generates a high current in the body. Several persistent musculoskeletal issues are treated with diathermy (10).

Recently, a new system has been created to transfer resistive and capacitive energy called TECAR therapy®. This type of diathermy increases the tissue's intrinsic metabolism through inter-tissue energy transfer without releasing energy to the outside. The TECAR device works based on the physical principles of the condenser and consists of two pads separated from the device, placed on the surface, and made of insulating material. One of the two electrodes, which is active, is in charge of transferring energy to the tissue. These two electrodes are connected to the energy-generating device, and a flow between them is established (10, 11). Effects of resistance-capacitance diathermy, also known as TECAR therapy, have increased microcirculation flow, vasodilation (increased oxygenation), and tissue heat (11, 12).

## 2. Objectives

Studies evaluating the efficacy of TECAR therapy in the treatment of peripheral neuropathies, particularly entrapments of peripheral nerves, are rarely available, although numerous studies have been conducted to date to investigate the effectiveness of TECAR therapy in various diseases. One of the pioneering studies on using TECAR therapy as an innovative physical therapy modality to treat median nerve entrapment at the wrist (carpal tunnel syndrome) was conducted in this study.

## 3. Methods

### 3.1. Trial Design

The present study is a single-blind randomized clinical trial performed in neuromuscular clinics affiliated with the Department of Physical Medicine and Rehabilitation of Isfahan University of Medical Sciences in 2022. Patients presenting with complaints in their hands, such as pain, paresthesia, numbness, or weakness, were approached with a detailed history and a complete physical examination, including special tests for CTS such as Compression, Tinel, Phalen, and Reverse Phalen tests. The patients were then chosen to receive an EMG-NCV neurophysiological evaluation to confirm the diagnosis of CTS and rule out any other potential diagnoses. Mild and moderate cases of CTS that met the criteria of the American Association of Electrodiagnostic Medicine's guidelines (13) were included in the study. A peak latency greater than 3.5 milliseconds and NCV across wrist < 40 m/s was considered mild, and an average motor onset latency greater than 4 milliseconds was considered moderate. The EMG-NCV was taken by the physical medicine specialist of Isfahan University of Medical Sciences with a Natus device (UltraPro S100, Natus Neurology Incorporated 3150 pleasant view road Middleton, WI USA).

Other inclusion criteria were age over 18 and under 60 years old and unilateral or bilateral idiopathic carpal tunnel syndrome lasting more than a month. The following criteria for exclusion were used: the presence of both systemic and local diseases (such as diabetes, RA, wrist arthritis, hypothyroidism), current or past cancer, use of a splint within the previous three months, pregnancy, burns to the hand or forearm, use of a pacemaker, prosthesis, or IUD, peripheral vascular diseases, fractures in the wrist and hand region, and corticosteroid injection in the carpal tunnel within the prior three months (14).

### 3.2. Interventions

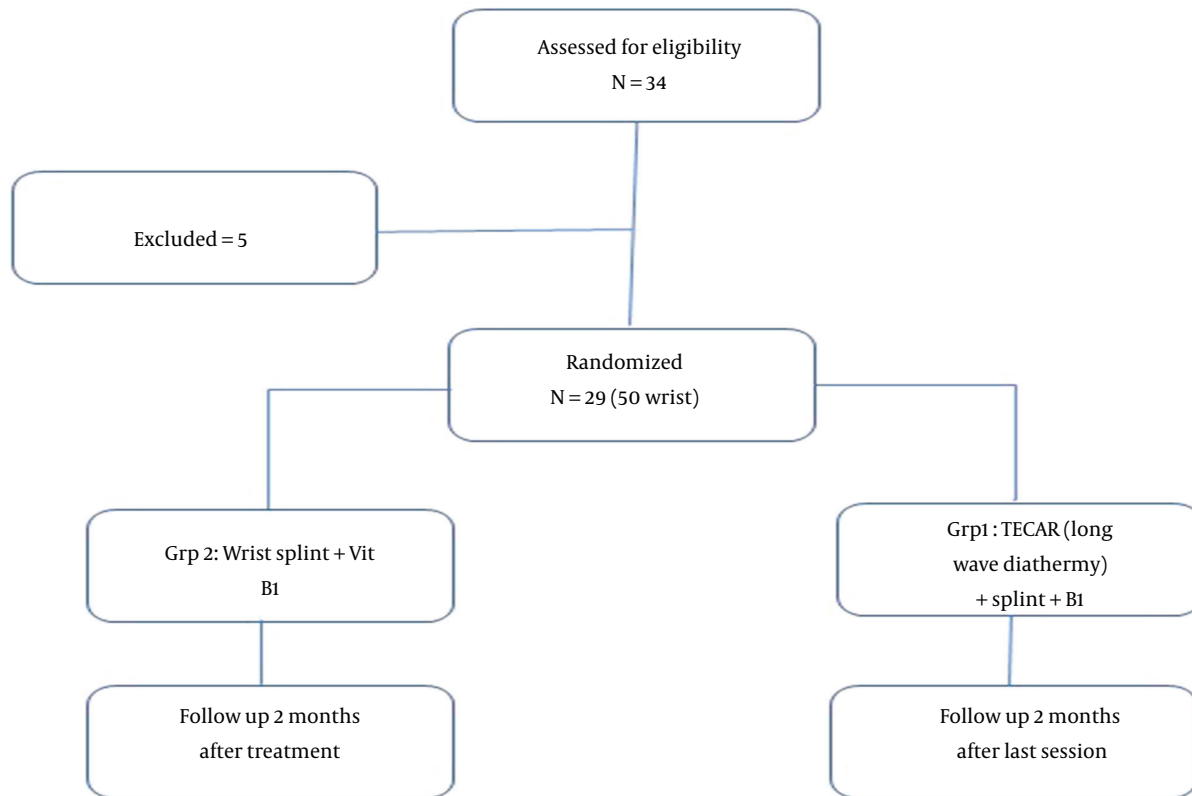
Both groups (1 and 2) were treated for four weeks with a wrist splint of thermoplastic type in 0 to 5 degrees of the extension during the nights and during activity in the daytime plus 300 mg of vitamin B1 (Vitamin B1-SHAF 300 MG) tablets. In the intervention group (group 1), TECAR therapy was also performed with the WINBACK 3 device made in France, with a frequency of 500 Hz and intensity of 30 to 50% and medium Capacitive Energy Transfer electrode (60 mm), from the proximal carpal tunnel to the palm Median nerve path, while the active capacitive electrode was moved by an operator in a circular pattern and the passive electrode fixed on the dorsum of the hand, in two sessions per week for four weeks (Figure 1).

### 3.3. Outcomes

At baseline, the VAS and Boston questionnaires were completed for both groups. After 2 months of the treatment, the questionnaires were filled out again. VAS scale (15) is a 10 cm graduated line whose numbers are graded from 0 (absence of pain) to 10 (the most severe pain possible). In this scale, the patient circles a number to determine its score. This scale has been widely and comprehensively used in research related to pain, and its validity and reliability have been confirmed (16).

Levine designed for the first time in 1993 the Boston Questionnaire to determine the severity and quality of carpal tunnel syndrome, which includes 2 sections on symptom severity and functional status in people with carpal tunnel syndrome (17). Each question has a maximum of five and a minimum of one point, and the average score is obtained by dividing the total points by the number of questions. Rezazadeh et al. (18) corroborated the validity and reliability of the Boston questionnaires among the Iranian population.

Clinical tests and NCV parameters (Median antidromic sensory peak latency, sensory nerve conduction velocity (NCV) across the wrist, Median motor distal onset latency) were also examined and compared before and after treatment.



**Figure 1.** Flowchart of the study

### 3.4. Sample Size

According to the formula for comparing means in clinical studies, considering type 1 error of .05, study power of 80%,  $d = 1.52$ , and  $S = 1.7$  (for VAS variable in previous studies (19)) for the interpretation of the result for the superiority of TECAR therapy combined to conventional treatment over conventional treatment, the final sample size was determined 25 hands in each group.

### 3.5. Randomization

The participants were enrolled using a consecutive sampling method. Random assignment was done via an online random sequence generator.

### 3.6. Blinding

In this single-blind research, the clinician and the patients were aware of the intervention groups. However, the outcome assessor and the statistical consultant were kept blind to each patient's intervention group.

### 3.7. Statistical Methods

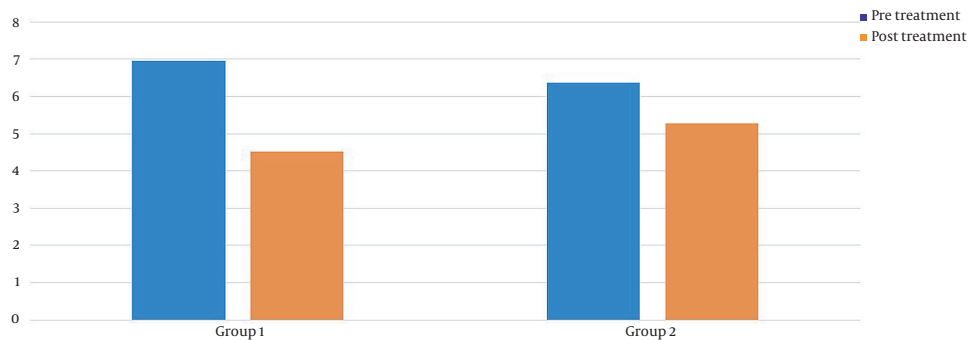
The collected data were analyzed using SPSS version 26. Appropriate parametric and nonparametric tests, including the paired-sample *t*-test (Boston-SSS and FSS variables and electrodiagnostic parameters in within-group comparison), independent *t*-test (Boston-SSS and FSS variables and electrodiagnostic parameters in between groups comparison), chi-square test (Demographic table variables), McNemar and Fisher exact test (Compression, Tinel, Phalen and Reverse Phalen variables within and between groups), Wilcoxon, and Mann-Whitney U test (VAS variable within and between groups), were used for statistical analyses.  $P$ -value  $< 0.05$  was considered as the significance level.

## 4. Results

Table 1 shows the demographics of the study participants. Before treatment, there was no significant difference in pain levels between the two groups. Two months after treatment, in both groups, patient pain levels were significantly reduced ( $P < 0.001$ ) (Figure 2). If the reduction of 25% of pain level (20) is considered significant, group 2

**Table 1.** Characteristics of the Study of Participants

	Group 1 (TECAR + Splint) (n = 14, 25 Wrist)	Group 2 (Splint) (n = 15, 25 Wrist)	P-Value
Age	47.08 ± 8.81	44.92 ± 10.20	0.678
Gender(f/m)	19/6	18/7	0.530
Affected wrist			0.321
Right	2	4	
Left	1	1	
Both	11	10	
Neurophysiological class			0.620
Mild	14	12	
Moderate	11	13	

**Figure 2.** VAS within groups

(wrist splint) didn't reach MCID (a minimal clinically important difference). There was a significant difference in pain reduction between the two groups, with a higher recovery rate in the TECAR treatment group ( $P = 0.008$ ) (Figure 3). After two months of treatment within groups, the difference became significant in all special clinical tests (Tinel (0.002), Compression (0.004), Phalen (0.031), and reverse Phalen (0.031)) in the TECAR group and Compression (0.031) and Reverse Phalen (0.016) tests in the wrist splint group. But between the groups, clinical tests did not show a significant difference. Before the study, there was no significant difference in BCTQ (FSS and SSS) variables between groups ( $P = 0.740$ ). Two months after the treatment, a significant difference was seen within ( $P = 0.018$ ,  $P = 0.01$ ) and between groups ( $P = 0.003$ ,  $P = 0.014$ ) (Table 2 and Figures 4-7). In our study, the wrist splint group didn't reach for both SSS and FSS to cut off level, but in the TECAR group, SSS was significant in MCID level (21), but FSS not significant.

Before the intervention, there was no discernible difference between the two groups according to neurophysiological tests (median sensory peak latency, median motor distal onset latency, and sensory NCV across the wrist). Also, there was no significant clinical difference between

and within the groups two months after the treatment, according to the tests (Table 3).

## 5. Discussion

In this study, we assessed the impact of TECAR therapy on electrodiagnostic parameters and clinical symptoms in patients with mild to moderate CTS.

In our study, VAS and Boston Questionnaire variables showed a good response, but NCV parameters did not show any specific changes from a clinical point of view. Perhaps the reason for the lack of clear changes in the NCS changes is that there is not enough time for changes to show. However, we can justify the VAS and Boston Questionnaire variables by examining the research and article review mechanism.

Few studies have been done on TECAR therapy, a novel deep heat physical modality, as a possible treatment for peripheral entrapment neuropathies or neuropathic pain.

The mechanism of action of TECAR has not been elucidated. The articles confirmed reduced muscle spasms, increased oxygenation, improved metabolic status, and reduced pain. By increasing radiofrequency energy to tis-

**Table 2.** Comparison of the Pre-treatment and Post-treatment Clinical Tests, Pain, Function, and Severity of Disease Within and Between Groups

	Group 1 (TECAR + Splint)	Group 2 (Splint)	95% Confidence Interval of the Difference		F	P-Value (Mann-Whitney)
			Lower	Upper		
<b>VAS</b>						
Pre	6.96 ± 1.01	6.36 ± 1.15	-0.01820	1.21820	0.0565	0.057
Post	4.52 ± .96	5.28 ± .97	-1.31235	-0.20765	0.019	0.008
P-value (Wilcoxon)	< 0.001	< 0.001				
						P-value (Independent t-test)
<b>Levine-Boston/SSS</b>						
Pre	2.30 ± .044	1.70 ± 0.48	3.628	9.492	0.111	0.740
Post	1.14 ± 0.15	1.45 ± 0.46	-5.571	-1.228	18.818	0.003
P-value (Paired-samples t-test)	0.012	0.011				
<b>Levine-Boston/FSS</b>						
Pre	1.95 ± 0.48	2.10 ± 0.44	-03.322	0.921	0.658	0.261
Post	1.45 ± 0.36	1.72 ± 0.38	-3.871	-0.448	< 0.001	0.014
P-value (Paired-samples t-test)	0.018	0.01				
						P-value (Fisher's Exact Test)
<b>Tinel test (positive/negative)</b>						
Pre	17/8	16/9				1.000
Post	7/18	12/13				0.378
P-value (McNemar test)	0.002	0.063				
<b>Compression test (positive/negative)</b>						
Pre	16/9	16/9				0.010
Post	8/17	10/15				
P-value (McNemar test)	0.004	0.031				0.194
<b>Phalen test (positive/negative)</b>						
Pre	13/12	15/10				0.015
Post	9/16	12/13				
P-value (McNemar test)	0.031	0.063				0.393
<b>Reverse Phalen test (positive/negative)</b>						
Pre	15/10	17/8				0.081
Post	9/16	10/15				0.667
P-value (McNemar test)	0.031	0.016				

sues, this treatment relaxes muscles and ligaments and improves nervous system repair (22). The main complaint of CTS patients is usually pain, which was measured by the VAS scale in this study. According to the Gate theory processes and mechanisms described above, reductions in VAS scales and patient pain levels are justifiable.

Lindblad et al. (23) investigated the efficacy of TECAR therapy for chemotherapy-induced neuropathic pain. Although TECAR therapy reduced patients' pain perception, the mean pain Numeric Rating Scale (NRS) did not reveal a discernible difference between groups. In the Lindblad

study, using the interferential therapy with high-power TECAR didn't result in significant pain relief in the intervention group; however, from the gate control theory, we expected some more pain reduction; this may be because the disease is chronic, and resistant to treatment, and different from a focal entrapment neuropathy. M. Nijalili et al. (24) evaluated the effectiveness of TECAR therapy in diabetic patients with peripheral neuropathy. Both groups (Group 1=TECAR + Infrared and Group 2=Infrared + Sham TECAR) experienced an improvement in pain management, with the intervention group experienc-

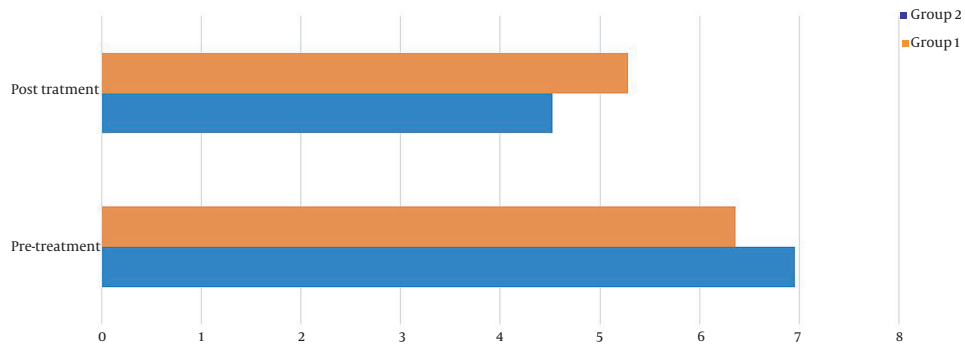


Figure 3. VAS- between groups

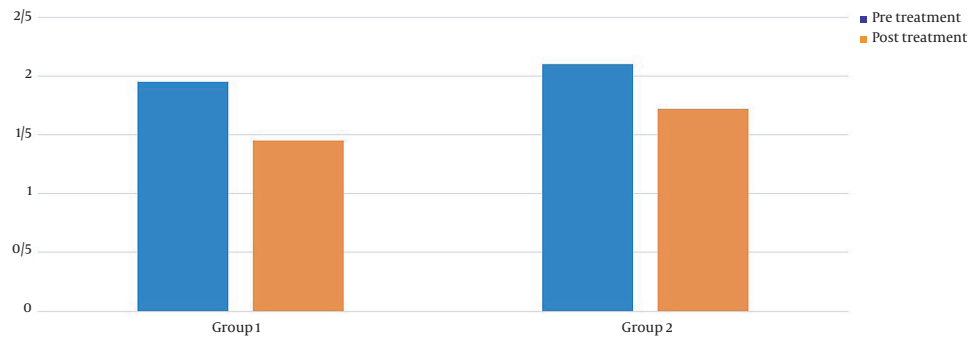


Figure 4. BCTQ- FSS- within groups

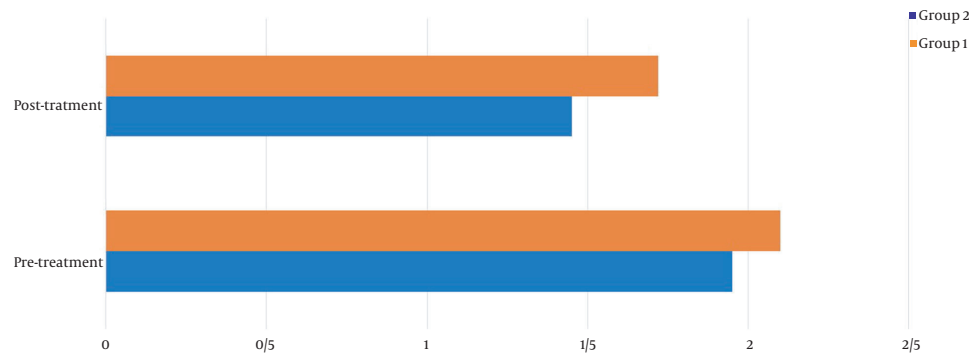


Figure 5. BCTQ- FSS- between groups

ing greater recovery. It suggests that the cause of the neuropathy may influence the TECAR therapy's effectiveness. In the field, however, more research is necessary.

Also, TECAR's action can be compared to other deep heat modalities. Ebenbichler et al. (25) studied the efficacy of therapeutic ultrasound in patients with mild to moderate CTS. They showed improved clinical symptoms, and

NCS parameters were also observed at the 6-month follow-up. Also, Huisstede et al. (26) reported no short-term improvements in their systematic review of ultrasound effects, but long-term effects were seen. We also found that TECAR therapy was effective in improving clinical symptoms but no NCS parameters at 2 months, during which long-term follow-up was not performed.

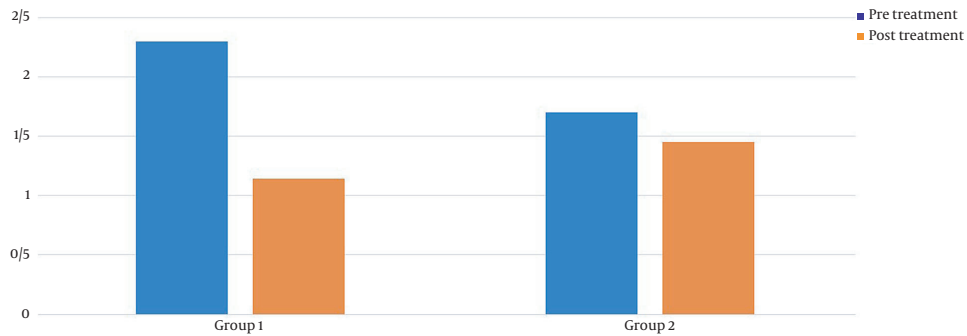


Figure 6. BCTQ-SSS-whin groups

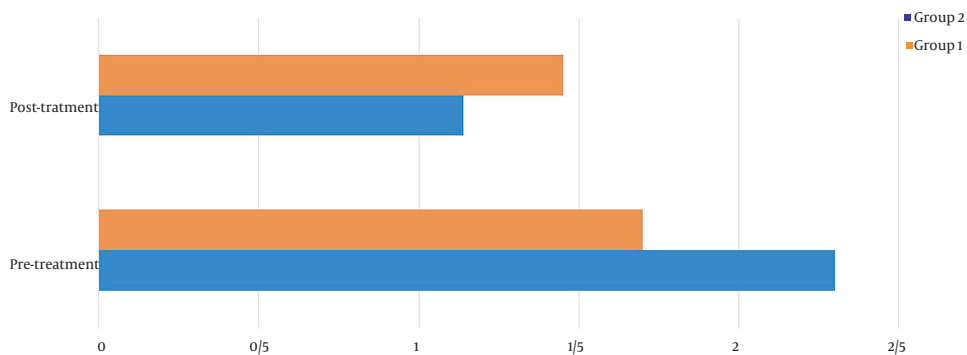


Figure 7. BCTQ-SSS-between groups

Possibly one of the reasons for reducing patient symptoms and improving performance is the reduction of edema in the carpal tunnel region. One of the pathological causes of this disease is increased pressure in this area, so by increasing lymphatic outflow and reducing pressure on nerves, gradual improvements in patient symptoms can be observed (27).

Another deep heat modality used for CTS treatment is shortwave diathermy (SWD). In a study by Incebiyik et al. (28), SWD has shown pain relief, improved hand function, and CTS clinical testing, but electrodiagnostic parameters have not been studied. In our study, the pain decreased, and hand function improved in the treatment group, but we observed no important changes in clinical tests compared to the control group.

Frasca et al. (6) conducted a study in 2011 to evaluate the efficacy of microwave diathermy (MWD) in sham and treatment groups. They found a reduction in VAS for pain and an improvement in the Boston Carpal Tunnel Questionnaire but no electrodiagnostic changes. In our study, an electrodiagnostic improvement was observed in addition to the improvements observed in the previous study.

Among other interventions in managing CTS, corticosteroid injections into the carpal tunnel revealed primary

evidence of efficacy in a systematic review by Piazzini et al. (29). Also, Visser et al. (30) showed that corticosteroid injection therapy is more effective in patients with mild CTS. In our study, sensory nerve conduction showed more changes than motor studies, but no significant differences were found between the two groups and clinically within the groups. This fact may indicate that patients with mild CTS respond better to treatment, but our study did not examine patients with mild and moderate CTS separately.

#### 5.1. Study Limitations

Because of COVID-19, the study was conducted with a small sample size, and no follow-up was performed due to time constraints. However, this study points to the need for longer follow-ups.

#### 5.2. Conclusions

Our results indicated that TECAR could effectively treat mild to moderate CTS patients. To investigate this, future interventional studies are required. It is better to conduct a study with a larger sample size and a longer follow-up to examine the changes in NCS from a clinical point of view.

**Table 3.** Comparison of Neurophysiologic Parameters Within and Between Groups

	Group 1 (TECAR+ Splint)	Group 2 (Splint)	95% Confidence Interval of the Difference		F	P-Value (Paired Sample t-Test)
			Lower	Upper		
<b>Median sensory PL (millisecond)</b>						
Pre	4.195 ± 0.322	4.170 ± 0.450	-0.197	0.248	2.179	0.818
Post	4.089 ± 0.284	4.130 ± 0.449	-0.256	-0.173	4.843	0.700
P-value (Independent t-test)	< 0.001	< 0.001				
<b>Median motor OL (millisecond)</b>						
Pre	4.209 ± 0.552	4.194 ± 0.448	-0.270	0.301	1.627	0.913
Post	4.147 ± 0.510	4.098 ± 0.400	-0.212	0.309	2.087	0.709
P-value (Independent t-test)	0.113	0.136				
<b>NCV across wrist (meter/second)</b>						
Pre	33.48 ± 5.94	34.72 ± 5.20	-4.414	1.942	0.221	0.438
Post	33.92 ± 5.99	35.12 ± 5.23	-4.825	1.545	0.237	0.306
P-value (Independent t-test)	0.048	< 0.001				

## Footnotes

**Authors' Contribution:** Study concept and design: Babak vahdatpour and Parisa Taheri.; analysis and interpretation of data: H.R.Ghasemi, and B.Vahdatpour.; drafting of the manuscript: F. M.; critical revision of the manuscript for important intellectual content: P.Taheri, and B.Vahdatpour.; statistical analysis: H.R.Ghasemi.

**Clinical Trial Registration Code:** This study was registered in the Iranian Registry of Clinical Trials (IRCT) by the code: [IRCT20220405054422N1](https://www.irct.ir/trial/20220405054422N1).

**Conflict of Interests:** The authors are not the editorial board management of this journal. Dr. Babak Vahdatpour is an interesting worker in the field of neuromuscular research and is a professor at Isfahan Medical University. The authors' other field is rehabilitation in the musculoskeletal system.

**Ethical Approval:** The Ethics Committee of Isfahan University of Medical Sciences approved the protocol of this study (code: [IR.MUI.MED.REC.1400.340](https://www.ir.mui.med.rec.1400.340)).

**Funding/Support:** There was no grant in this study.

**Informed Consent:** The participants were aware of the study goals and protocol. Written consent was obtained from patients before assignment to each treatment group.

## References

- Tadgerbashi K, Akesson A, Atroushi I. Incidence of referred carpal tunnel syndrome and carpal tunnel release surgery in the general population: Increase over time and regional variations. *J Orthop Surg (Hong Kong)*. 2019;**27**(1):2309499019825570. [PubMed ID: [30798784](https://pubmed.ncbi.nlm.nih.gov/30798784/)]. <https://doi.org/10.1177/2309499019825572>.
- Luchetti R, Tognon S, Cacciavillani M, Ronco S, Buzzelli N, Lanni G. Observational multicentric survey on carpal tunnel syndrome: demographic and clinical data from 34 Italian centers. *Eur Rev Med Pharmacol Sci*. 2017;**21**(3):460-9. [PubMed ID: [28239826](https://pubmed.ncbi.nlm.nih.gov/28239826/)].
- Stevens JC, Smith BE, Weaver AL, Bosch EP, Deen HJ, Wilkens JA. Symptoms of 100 patients with electromyographically verified carpal tunnel syndrome. *Muscle Nerve*. 1999;**22**(10):1448-56. [PubMed ID: [10487914](https://pubmed.ncbi.nlm.nih.gov/10487914/)]. [https://doi.org/10.1002/\(sici\)1097-4598\(199910\)22:10<1448::aid-must17>3.0.co;2-y](https://doi.org/10.1002/(sici)1097-4598(199910)22:10<1448::aid-must17>3.0.co;2-y).
- Bobowik PZ. Effectiveness of physiotherapy in carpal tunnel syndrome (CTS). *Adv Rehabil*. 2019;**2019**(2):47-58. <https://doi.org/10.5114/areh.2019.85023>.
- Michelsen H, Posner MA. Medical history of carpal tunnel syndrome. *Hand Clin*. 2002;**18**(2):257-68. [PubMed ID: [12371028](https://pubmed.ncbi.nlm.nih.gov/12371028/)]. [https://doi.org/10.1016/s0749-0712\(01\)00006-3](https://doi.org/10.1016/s0749-0712(01)00006-3).
- Frasca G, Maggi L, Padua L, Ferrara PE, Granata G, Minciotti I, et al. Short-term effects of local microwave hyperthermia on pain and function in patients with mild to moderate carpal tunnel syndrome: a double blind randomized sham-controlled trial. *Clin Rehabil*. 2011;**25**(12):1109-18. [PubMed ID: [21937521](https://pubmed.ncbi.nlm.nih.gov/21937521/)]. <https://doi.org/10.1177/0269215511400767>.
- Padua L, Coraci D, Erra C, Pazzaglia C, Paolasso I, Loreti C, et al. Carpal tunnel syndrome: clinical features, diagnosis, and management. *Lancet Neurol*. 2016;**15**(12):1273-84. [PubMed ID: [27751557](https://pubmed.ncbi.nlm.nih.gov/27751557/)]. [https://doi.org/10.1016/S1474-4422\(16\)30231-9](https://doi.org/10.1016/S1474-4422(16)30231-9).
- Padua L, Padua R, Lo Monaco M, Aprile I, Tonali P. Multiperspective assessment of carpal tunnel syndrome: a multicenter study. Italian CTS Study Group. *Neurology*. 1999;**53**(8):1654-9. [PubMed ID: [10563608](https://pubmed.ncbi.nlm.nih.gov/10563608/)]. <https://doi.org/10.1212/wnl.53.8.1654>.
- Huisstede BM, Friden J, Coert JH, Hoogvliet P, European HG. Carpal tunnel syndrome: hand surgeons, hand therapists, and physical medicine and rehabilitation physicians agree on a multidisciplinary treatment guideline-results from the European HANDGUIDE Study. *Arch Phys Med Rehabil*. 2014;**95**(12):2253-63. [PubMed ID: [25127999](https://pubmed.ncbi.nlm.nih.gov/25127999/)]. <https://doi.org/10.1016/j.apmr.2014.06.022>.
- Hawamdeh M. The effectiveness of capacitive resistive diathermy (Tecartherapy®) in acute and chronic musculoskeletal lesions and pathologies. *Eur J Sci Res*. 2014;**118**(3):336-40.
- Ganzit GP, Stefanini L, Stesina G. Tecar® Therapy in the Treatment of Acute and Chronic Pathologies In Sports. *FMSI Ital Sports Med Fed CONI Ins Sports Med*. 2000.
- Onesta E. Hyperthermia through Resistive and Capacitive Energy Transfer in the Treatment of Acute and Chronic Musculoskeletal Lesions. *La Riabilitazione*. 1998.



13. Werner RA, Andary M. Electrodiagnostic evaluation of carpal tunnel syndrome. *Muscle Nerve*. 2011;**44**(4):597-607. [PubMed ID: 21922474]. <https://doi.org/10.1002/mus.22208>.
14. Bilgici A, Ulusoy H, Kuru O, Canturk F. The comparison of ultrasound treatment and local steroid injection plus splinting in the carpal tunnel syndrome: a randomized controlled trial. *Bratisl Lek Listy*. 2010;**111**(12):659-65. [PubMed ID: 21384736].
15. McGrath PJ, Finley GA. Measurement of pain. *Annales Nestle*. Nestec Ltd; 1999. p. 13-9.
16. Zahednezhad SH, Salehi R, Tajali SH, Borji A. Correlation between pain intensity and disability level with some of the impairments in patients with nonspecific low back pain. *J Ilam Univ Med Sci*. 2013;**21**(2):10-20.
17. Fischer J, Thompson NW, Harrison JW. A Self-Administered Questionnaire for the Assessment of Severity of Symptoms and Functional Status in Carpal Tunnel Syndrome. *Classic Papers in Orthopaedics*. 2014. p. 349-51.
18. Rezazadeh A, Bakhtiari AH, Samaei A, Moghimi J. Validity and reliability of the Persian Boston questionnaire in Iranian patients with carpal tunnel syndrome. *Koomesh*. 2014;**15**(2):138-45.
19. Paolucci T, Pezzi L, Centra MA, Porreca A, Barbato C, Bellomo RG, et al. Effects of capacitive and resistive electric transfer therapy in patients with painful shoulder impingement syndrome: a comparative study. *J Int Med Res*. 2020;**48**(2):300060519883090. [PubMed ID: 31680597]. [PubMed Central ID: PMC7783264]. <https://doi.org/10.1177/0300060519883090>.
20. Hsu PC, Liao KK, Lin KP, Chiu JW, Wu PY, Chou CL, et al. Comparison of Corticosteroid Injection Dosages in Mild to Moderate Idiopathic Carpal Tunnel Syndrome: A Randomized Controlled Trial. *Arch Phys Med Rehabil*. 2020;**101**(11):1857-64. [PubMed ID: 32682938]. <https://doi.org/10.1016/j.apmr.2020.06.018>.
21. Ozer K, Malay S, Toker S, Chung KC. Minimal clinically important difference of carpal tunnel release in diabetic and nondiabetic patients. *Plast Reconstr Surg*. 2013;**131**(6):1279-85. [PubMed ID: 23416439]. [PubMed Central ID: PMC4787587]. <https://doi.org/10.1097/PRS.0b013e31828bd6ec>.
22. Oh D, Kim S, Yoo K. Effect of Physiotherapeutic Intervention Using TECAR Therapy on Pain Self-Awareness and Hip Joint Function in Hip Impingement Syndrome: A Case Study. *J Korean Soc Phys Med*. 2021;**16**(3):45-53. <https://doi.org/10.13066/kspm.2021.16.3.45>.
23. Lindblad K, Bergkvist L, Johansson AC. Evaluation of the treatment of chronic chemotherapy-induced peripheral neuropathy using long-wave diathermy and interferential currents: a randomized controlled trial. *Support Care Cancer*. 2016;**24**(6):2523-31. [PubMed ID: 26687020]. <https://doi.org/10.1007/s00520-015-3060-7>.
24. Niajalili M, Sedaghat M, Rezasoltani A, Akbarzade Baghban AR, Naimi SS. Effect of Capacitive Tecar Therapy on Foot Pain and Tactile Sensation in Patients with Type 2 Diabetes. *J Rehabil*. 2020;**21**(3):304-19. <https://doi.org/10.32598/rj.21.3.60.5>.
25. 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;**316**(7133):731-5. [PubMed ID: 9529407]. [PubMed Central ID: PMC28476]. <https://doi.org/10.1136/bmj.316.7133.731>.
26. Huisstede BM, Hoogvliet P, Randsdorp MS, Glerum S, van Middekoop M, Koes BW. Carpal tunnel syndrome. Part I: effectiveness of nonsurgical treatments—a systematic review. *Arch Phys Med Rehabil*. 2010;**91**(7):981-1004. [PubMed ID: 20599038]. <https://doi.org/10.1016/j.apmr.2010.03.022>.
27. Cau N, Cimolin V, Aspesi V, Galli M, Postiglione F, Todisco A, et al. Preliminary evidence of effectiveness of TECAR in lymphedema. *Lymphology*. 2019;**52**(1):35-43. [PubMed ID: 3119913].
28. Incebiyik S, Boyaci A, Tutoglu A. Short-term effectiveness of short-wave diathermy treatment on pain, clinical symptoms, and hand function in patients with mild or moderate idiopathic carpal tunnel syndrome. *J Back Musculoskeletal Rehabil*. 2015;**28**(2):221-8. [PubMed ID: 25061038]. <https://doi.org/10.3233/BMR-140507>.
29. Piazzini DB, Aprile I, Ferrara PE, Bertolini C, Tonali P, Maggi L, et al. A systematic review of conservative treatment of carpal tunnel syndrome. *Clin Rehabil*. 2007;**21**(4):299-314. [PubMed ID: 17613571]. <https://doi.org/10.1177/0269215507077294>.
30. Visser LH, Ngo Q, Groeneweg SJ, Brekelmans G. Long term effect of local corticosteroid injection for carpal tunnel syndrome: a relation with electrodiagnostic severity. *Clin Neurophysiol*. 2012;**123**(4):838-41. [PubMed ID: 21962473]. <https://doi.org/10.1016/j.clinph.2011.08.022>.