Efficacy of splinting and oral steroids in the treatment of carpal tunnel syndrome: A prospective randomized clinical and electrophysiological study

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Objective: To study the efficacy of splinting and oral steroids in the management of carpal tunnel syndrome (CTS).

Design: Prospective, randomized, open-label, clinical and electrophysiological study with 3-month follow-up.

Materials and Methods: Forty patients with CTS were randomly divided into splint group (N-20), wearing splint in neutral position for 4 weeks; and steroid group (N-20), who received oral prednisolone 20 mg/day for 2 weeks followed by 10 mg/day for 2 weeks. Clinical and electrophysiological evaluations were done at baseline and at 1-month and 3-month follow-up. Independent 't' test and paired 't' test were used for statistical analysis. Outcome Measures: Primary outcome measure was the symptom severity score and functional status score. Secondary outcome measures were median nerve sensory and motor distal latency and conduction velocity. Results: At the end of 3 months, statistically significant improvement was seen in symptom severity score and functional status score in both groups (P<0.001). Median nerve sensory distal latency and conduction velocity also improved significantly in both the groups at 3 months. Improvement in motor distal latency was significant (P=0.001) at 3 months in steroid group, while insignificant improvement (P=0.139) was observed in splint group. On comparing the clinical and electrophysiological improvement between the two groups, except for the functional status score, there was no significant difference at 3-month follow-up. Improvement in functional status score was significantly more in steroid group (P=0.03). Conclusion: There was significant improvement in both groups, clinically as well as electrophysiologically, at 3 months. On comparing the efficacy of the two treatment methods, except for the functional status score, there was no significant difference between the two groups.

Key words: Carpal tunnel syndrome, clinical follow-up, electrophysiology, oral steroids, splinting

Introduction

Carpal tunnel syndrome (CTS) is the most common focal compressive neuropathy and is caused by entrapment of the median nerve in the carpal tunnel. Prevalence of electrophysiologically confirmed symptomatic CTS is about 3% among women and 2% among men, with peak prevalence in women older than 55 years of age. Early diagnosis and treatment of CTS are important because delay can result in irreversible median nerve damage with persistent symptoms and permanent disability. Treatment options include rest and avoidance of excessive activity at hand, exercises, laser therapy, splinting the wrist in neutral position, nonsteroidal anti-inflammatory drugs oral steroids, local corticosteroid injections and surgery to decompress the median nerve segment. Splinting the wrist in neutral position has been reported to provide symptomatic relief. Oral steroid therapy has also been found to result in significant decrease in symptoms of CTS in a few recent studies. However, data on comparative analysis of the conservative treatment modalities for CTS is limited in the literature. The present study aims at studying the effects of splinting in neutral position of the wrist and oral prednisolone on symptomatology and neurophysiological variables in patients with CTS.

Materials and Methods

A prospective randomized open-label clinical and electrophysiological study of efficacy of splinting and oral steroids for the treatment of CTS was done. Study population consisted of 66 patients who were diagnosed to have CTS in neurology outpatient department of a tertiary care center between July 2003 and December 2004. The study was approved by the ethical research committee of the Institute. Patients with symptoms suggestive of carpal tunnel syndrome of at least 1-month duration and electrophysiological evidence of median neuropathy at wrist were included in the study. Patients...
were referred from internal medicine, plastic surgery and orthopedic outpatient departments of our institute and also by general practitioners from outside the hospital. The clinical criteria laid down by the American Academy of Neurology were used for diagnosis of CTS.\textsuperscript{[8]} The electrophysiological criteria used for the diagnosis of CTS included the presence of two or more of the following:

1. Median nerve motor distal latency recording at abductor pollicis brevis and stimulating at wrist greater than 4.4 ms.
2. Median nerve antidromic sensory peak latency recording at digit II greater than 3.5 ms.
3. Difference between antidromic median sensory latency and ulnar sensory latency at digit IV greater than 0.5 ms.
4. Antidromic latency difference of $>0.5$ ms between median nerve at digit II and ulnar nerve at digit V using the same distance of measurement.

The severity of symptoms and functional status was assessed using a scoring system as proposed by Levine \textit{et al.}\textsuperscript{[9]} Symptom severity score has 11 items concerning pain, nocturnal symptoms, numbness, tingling and weakness. Functional status score has 8 items (difficulty in writing, buttoning, opening jars, holding a book, gripping telephone handle, household chores, carrying grocery bags, bathing and dressing). Each item of these scales has five responses ranging from 1 (no symptom or no difficulty) to 5 (very severe symptoms preventing the activity). Mean of the scores for all symptoms and activities was calculated and compared with scores at follow-up at 1 month and 3 months. All patients were randomly allocated to one of the following two groups:

1. Splinting in neutral position.
2. Oral steroid.

Randomization was done using the table of random numbers. Patients in splint group were advised to wear commercially available carpal tunnel splint at night and as much as possible during the daytime for 4 weeks. In the case of bilateral symptoms, both hands were treated. They were also told not use additional medicines or other methods of treatment during the study period.

Patients in steroid group were given oral prednisolone 20 mg/day for 2 weeks followed by 10 mg/day for another 2 weeks, as described in previous studies.\textsuperscript{[9]} Advice to avoid extremes of wrist flexion or extension, excessive hand movement and hand rest was common to both groups. The clinical outcome was assessed at 1 and 3 months. Primary outcome measures were the symptom severity score and functional status score. Secondary outcome measures were median nerve sensory and motor distal latency and conduction velocity. Normal values of the motor conduction velocity and sensory conduction velocity in our lab are 56 ± 10 m/s and 55 ± 12 m/s respectively. Nerve conduction study was performed at baseline and also at 1- and 3-month follow-up.

\textbf{Exclusion criteria}

1. Patient with diabetes mellitus, trauma to wrist and deformity.
3. Patients with advanced CTS having wasting, marked weakness with marked axonal loss on nerve conduction study or nonstimulatable nerves.
4. Patients with a history of peptic ulcer.
5. Patients treated previously for CTS using medical or surgical therapy.
6. Pregnant women with CTS.
7. Patients with systemic disorders like rheumatoid arthritis, hypothyroidism, amyloidosis, etc.

\textit{Nerve conduction study was performed on Medtronic machine using Keypoint software version 3.21. Motor and sensory conduction studies of median and ulnar nerve were performed on both hands. Motor conduction for median nerve were performed using the surface electrodes over abductor pollicis brevis (active) and first metacarpophalangeal joint (reference) while stimulation was done at wrist and elbow. For ulnar motor conduction, active electrode was placed at abductor digiti minimi and reference at the fifth metacarpophalangeal joint while stimulation was done at wrist, below elbow and above elbow. Antidromic sensory nerve conduction was performed using standard ring electrode recordings from digit II for median and digit V for ulnar nerve. The recordings were made from ring finger with median and ulnar stimulation at wrist for digit IV sensory study. Parameters including distal latency, amplitude, conduction velocity and F responses were recorded.}

All statistical calculations were performed using statistical package for social sciences (SPSS) for window version 10. Unpaired ‘t’ test was used to calculate statistical significance when comparing baseline parameters, improvement in clinical scores and change in electrophysiological parameters between the two groups. Chi-square test was used to compare sex distribution between the two groups at baseline. For comparison of variables within the groups between baseline and 1 month and 3 months post-treatment, paired ‘t’ test was used. \textit{P} value of less than 0.05 was considered significant.

\textbf{Results}

This study included 66 patients with 117 CTS-affected hands. Of the 66 patients, 26 did not fulfill the inclusion criteria and therefore were excluded. Of these, 13 patients had CTS secondary to conditions like diabetes mellitus,\textsuperscript{[2]} hypothyroidism,\textsuperscript{[6]} pregnancy,\textsuperscript{[5]} rheumatoid arthritis,\textsuperscript{[1]} amyloidosis.\textsuperscript{[1]} Remaining 13 patients had gross wasting, markedly decreased SNAP/CMAP amplitude or unrecordable SNAP/CMAPs in median nerve.

Remaining 40 patients (with 71 affected hands) were randomly allocated to two groups. Splint group had 20 patients with 36 affected hands and steroid group had 20 patients with 35 affected hands. Baseline characteristics of these patients are shown in Table 1. Mean age of the patients in the splint group was 42.91 years with a range of 23 to 60 years, while in steroid group it was 41.57 years with a range of 28 to 60 years [Table 1]. There was statistically no significant difference between the age and sex distribution of the patients in two groups. Overall, incidence of CTS was almost five times more in females compared to males. Electrophysiological abnormalities in both the groups at baseline.
are shown in Table 2. There were no statistically significant differences between the two groups.

Tables 3 and 4 show the outcome of treatment in both the treatment groups. In splint group, there was significant (P-value <0.001) improvement in symptom severity score, functional status score and sensory distal latency at 1-month follow-up and the same was sustained at 3-month follow-up. Sensory conduction velocity also showed statistically significant improvement at 1- and 3-month follow-up. In steroid group also, there was significant improvement in symptom severity score, functional status score at 1-month follow-up and the same was sustained at 3-month follow-up. Sensory and motor distal latencies and sensory conduction velocity showed statistically significant improvement at 3-month follow-up. Motor conduction velocity did not show significant improvement at 1- or 3-month follow-up.

Differences of the mean between baseline and 1 month and 3 month were compared between the two groups (Table 5). Functional status score showed significant difference at 1 month and 3 month (better results in steroid group). Sensory conduction velocity also showed significant difference at 1 month, (more improvement in steroid group), but at 3 months the change from baseline was not significant between the two groups. Rest of the parameters including symptom severity score, motor distal latency, motor conduction velocity and sensory distal latency did not show any significant difference either at 1- or 3-month follow-up. Two patients in steroid group and one in splint group had worsening of symptoms at 3-month follow-up after initial response at 1 month.

Overall, significant improvement in primary outcome measures was observed at 1- and 3-month follow-up in both the groups and most of the electrophysiological data (secondary outcome

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Splint group (N-20 with 36 hands)</th>
<th>Steroid group (N-20 with 35 hands)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>42.91 ± 9.39</td>
<td>41.57 ± 9.26</td>
<td>0.54</td>
</tr>
<tr>
<td>(mean ± SD)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sex (male/female)</td>
<td>3/17</td>
<td>4/16</td>
<td>0.67</td>
</tr>
<tr>
<td>BMI (mean ± SD)</td>
<td>23.87 ± 2.23</td>
<td>23.84 ± 2.41</td>
<td>0.95</td>
</tr>
<tr>
<td>Duration (months)</td>
<td>6.40 ± 7.09</td>
<td>6.31 ± 7.50</td>
<td>0.95</td>
</tr>
</tbody>
</table>

FSS: Functional status score, SSS: Symptom severity score. P-value <0.05 is significant.

### Table 2: Electrophysiological abnormalities at baseline

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Splint group (36 hands) (mean ± SD)</th>
<th>Steroid group (36 hands) (mean ± SD)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDL (milliseconds)</td>
<td>3.94 ± 0.56</td>
<td>3.94 ± 0.98</td>
<td>0.982</td>
</tr>
<tr>
<td>MCV (meters/second)</td>
<td>54.13 ± 14.65</td>
<td>53.20 ± 8.72</td>
<td>0.749</td>
</tr>
<tr>
<td>SNAP amplitude (microvolts)</td>
<td>22.86 ± 9.70</td>
<td>26.40 ± 9.36</td>
<td>0.123</td>
</tr>
<tr>
<td>SCV (meters/second)</td>
<td>33.04 ± 8.69</td>
<td>36.31 ± 8.82</td>
<td>0.121</td>
</tr>
</tbody>
</table>

MDL: Motor distal latency, MCV: Motor conduction velocity, CMAP: Compound muscle action potential, SDL: Sensory distal latency, SCV: Sensory conduction velocity, SNAP: Sensory nerve action potential, P-value<0.05 is significant.

### Table 3: Treatment outcome: Splint group (N-36)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Baseline</th>
<th>1 month</th>
<th>3 months</th>
<th>Mean change (0-1) mean ± SD (P value)</th>
<th>Mean change (0-3) mean ± SD (P value)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SSS</td>
<td>2.83 ± 0.41</td>
<td>2.48 ± 0.52</td>
<td>2.43 ± 0.56</td>
<td>0.54 ± 0.42 (&lt;0.001)</td>
<td>0.39 ± 0.54 (&lt;0.001)</td>
</tr>
<tr>
<td>FSS</td>
<td>1.74 ± 0.34</td>
<td>1.60 ± 0.41</td>
<td>1.57 ± 0.43</td>
<td>0.14 ± 0.13 (&lt;0.001)</td>
<td>0.16 ± 0.17 (&lt;0.001)</td>
</tr>
<tr>
<td>MDL</td>
<td>3.94 ± 0.56</td>
<td>3.88 ± 0.66</td>
<td>3.74 ± 0.51</td>
<td>0.06 ± 0.77 (0.607)</td>
<td>0.20 ± 0.79 (0.139)</td>
</tr>
<tr>
<td>MCV</td>
<td>54.13 ± 14.65</td>
<td>52.03 ± 9.42</td>
<td>50.73 ± 7.46</td>
<td>2.09 ± 16.53 (0.452)</td>
<td>3.40 ± 16.22 (0.217)</td>
</tr>
<tr>
<td>SDL</td>
<td>4.23 ± 0.83</td>
<td>4.07 ± 0.86</td>
<td>3.88 ± 0.56</td>
<td>0.16 ± 0.63 (0.136)</td>
<td>0.35 ± 0.76 (0.009)</td>
</tr>
<tr>
<td>SCV</td>
<td>33.04 ± 8.69</td>
<td>34.40 ± 9.51</td>
<td>35.87 ± 9.06</td>
<td>-1.41 ± 7.11 (0.24)</td>
<td>-2.83 ± 7.88 (0.03)</td>
</tr>
</tbody>
</table>

SSS: Symptom severity score. SCV: Sensory conduction velocity, FSS: Functional status score, MCV: Motor conduction velocity, SDL: Sensory distal latency, MDL: Motor distal latency. 0-1: Difference of mean between baseline and 1 month follow-up, 0-3: Difference of mean between baseline and 3 months follow-up, P-value<0.05 is significant.

### Table 4: Treatment outcome: Steroid group (N-35)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Baseline</th>
<th>1 month</th>
<th>3 months</th>
<th>Mean change (0-1) mean ± SD (P value)</th>
<th>Mean change (0-3) mean ± SD (P value)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SSS</td>
<td>2.67 ± 0.37</td>
<td>2.27 ± 0.48</td>
<td>2.18 ± 0.63</td>
<td>0.40 ± 0.30 (&lt;0.001)</td>
<td>0.49 ± 0.44 (&lt;0.001)</td>
</tr>
<tr>
<td>FSS</td>
<td>1.68 ± 0.31</td>
<td>1.48 ± 0.30</td>
<td>1.45 ± 0.35</td>
<td>0.20 ± 0.11 (&lt;0.001)</td>
<td>0.23 ± 0.17 (&lt;0.001)</td>
</tr>
<tr>
<td>MDL</td>
<td>3.94 ± 0.98</td>
<td>3.74 ± 0.73</td>
<td>3.47 ± 0.77</td>
<td>0.19 ± 0.76 (0.143)</td>
<td>0.47 ± 0.73 (0.001)</td>
</tr>
<tr>
<td>MCV</td>
<td>53.20 ± 8.72</td>
<td>52.17 ± 6.19</td>
<td>54.01 ± 5.64</td>
<td>1.03 ± 11.37 (0.594)</td>
<td>-0.80 ± 8.72 (0.588)</td>
</tr>
<tr>
<td>SDL</td>
<td>3.88 ± 1.02</td>
<td>3.74 ± 0.80</td>
<td>3.32 ± 0.64</td>
<td>0.13 ± 0.71 (0.27)</td>
<td>0.55 ± 0.66 (&lt;0.001)</td>
</tr>
<tr>
<td>SCV</td>
<td>36.31 ± 8.82</td>
<td>33.75 ± 5.89</td>
<td>39.82 ± 6.47</td>
<td>2.56 ± 9.38 (0.11)</td>
<td>-3.51 ± 8.48 (0.02)</td>
</tr>
</tbody>
</table>

SSS - Symptom severity score, SCV: Sensory conduction velocity, FSS - Functional status score, MCV-Motor conduction velocity, SDL: Sensory distal latency, MDL: Motor distal latency. 0-1: Difference of mean between baseline and 1 month follow-up, 0-3: Difference of mean between baseline and 3 months follow-up, P-value<0.05 is significant.
In the splint group, there was significant improvement in the symptom severity scores and functional status scores at 1 month and the same was sustained at 3-month follow-up. Along with symptomatic improvement, there was significant improvement in sensory distal latency and sensory conduction velocity during the 3-month follow-up. However, there was no significant change in motor nerve conduction parameters, possibly because the motor conduction abnormalities itself were minimal at baseline. In the published literature, there are variable reports of success with splint use (31 to 67%) in different studies.\(^\text{[4,10]}\) Gerristen et al reported a success rate of 31% (26 of 83 patients) with splinting.\(^\text{[10]}\)

Patients in steroid group also showed significant improvement in clinical scores at 1- and 3-month follow-up. Sensory and motor distal latencies and sensory conduction velocities also showed significant improvement at 3-month follow-up. In a study by Herskovitz et al,\(^\text{[7]}\) all the treated patients reported improvement at 2-week follow-up of treatment with oral steroids. However, this effect gradually waned off over 8 weeks in all of them. Chang et al\(^\text{[6]}\) compared a regimen of 20 mg prednisolone per day for 2 weeks followed by 10 mg daily for next 2 weeks, with 20 mg per day of prednisolone for two weeks only. Improvement in Global Symptom Score in the 4-week treatment group was achieved in 66% of patients. In the 2-week treatment group, the improvement was seen in 49% of patients. The difference was statistically not significant. Persistence of improvement was 74.20% in the 4-week group and 74.10% in the 2-week group with no significant difference. At 1-year follow-up, improvement decreased to 51%, suggesting relapse of symptoms in 17% of patients who had responded at 1 month.

On comparing the mean of the change from baseline to 1 and 3 month (0-1 and 0-3) between the two groups, there was no significant difference in symptom severity score, motor and sensory distal latencies and conduction velocities either at 1-or 3-month follow-up. However, functional status score showed significantly more improvement in steroid group at 3-month follow-up. To the best of our knowledge, there are no comparative studies of these two treatment methods in the published literature for comparison with our data. Based on these observations, we do not intend to propose that oral steroid therapy is the treatment of choice for CTS or that splinting is inferior to steroids. However, findings do suggest the need for further studies on this subject. While deciding about the best approach for treatment, one would also have to consider the adverse effects like Cushingoid features, avascular necrosis, gastrointestinal side effects, worsening of diabetes and hypertension, etc., which might develop due to repeated use of systemic steroids since the symptoms of CTS may be recurrent.

Two patients in the steroid group and one in the splint group had worsening of symptoms at 3-month follow-up after initial response at 1 month. In the study by Burke et al,\(^\text{[4]}\) 17% of the patients reported worsening of symptoms between the 2-week and 2-month follow-up after splint use. Chang et al\(^\text{[6]}\) also reported that at 1-year follow-up, 17% of patients who had responded at 1 month had a relapse of symptoms at 1-year.

### Discussion

This was a prospective study of the efficacy of two treatment modalities—splint and oral steroids—in patients with CTS. We chose oral steroids since steroid injections are known to cause mechanical or chemical nerve injury and several patients do not accept injections in the wrist. Age and sex distribution in this study is similar to what has been observed by studies from India and other countries.\(^\text{[13,16]}\) Overall positivity of tinel sign and phalen sign was 38 and 62% respectively. Positivity of tinel sign has been reported to vary from 9 to 73% and that of phalen sign varies from 10 to 73%.\(^\text{[15]}\) Murthy et al reported a positivity of 24.6% for tinel sign and 17.5% for phalen sign.\(^\text{[16]}\) The most common abnormality on electrodiagnostic study was sensory distal latency prolongation. Of 71 CTS-affected hands, 5.63% had normal absolute values of latencies, amplitude and conduction velocities, but the difference between median and ulnar sensory distal latencies was consistent with the diagnosis of CTS.\(^\text{[14]}\) Electrodiagnosis of CTS is likely to be missed in 10 to 25% of patients if such comparison studies are not performed.\(^\text{[15]}\)
follow-up. In fact, all the conservative methods of treatment for CTS do not have a sustained effect. In a recent study, splinting was combined with steroid injections and patients were followed up for 1 year. The study concluded that steroid injections and splinting are effective for relief of CTS symptoms but have a long-term effect in only 10% of patients.\cite{17} In the study by Gelberman et al.,\cite{89} 89% of the patients with severe CTS (having constant numbness, weakness, atrophy or sensory loss) had recurrence within a year, while only 60% patients with mild CTS (intermittent symptoms and normal sensory and motor examination) had recurrence after splinting and local injection, suggesting that relatively advanced CTS is more likely to recur after conservative treatments.\cite{18}

The obvious limitations of this study are short follow-up and small number of subjects in both the groups. However, our study has some methodological superiority also – like prospective, randomized design and the use of well established validated clinical scores for clinical assessment. We do not consider placebo effect as a possibility for the improvement in our study since both clinical and electrodiagnostic measurements improved at the 3-month follow-up. Nonetheless, to be very sure, randomized, double blind, placebo-controlled study with long-term follow-up is needed.

### Conclusions

Both the treatment methods (splint and oral steroids) were found to be effective. There was significant improvement in both groups clinically during follow-up at 1 and 3 months as well as electrophysiologically at 3 months. On comparing the efficacy of these two treatment methods, there was no significant difference at 3-month follow-up except for the functional status score. Improvement in functional status score was significantly more in steroid group. Oral steroids are safe and well tolerated at low dose in the short term. Studies with larger sample size and longer follow-up are suggested to establish the duration of beneficial effect, superiority of one treatment method over the other and actual relapse rates of symptoms.

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### References


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