

DISSERTATION

ON

"EVALUATION OF CLINICAL, ELECTROPHYSIOLOGICAL, AND RADIOLOGICAL PROFILE
OF CARPAL TUNNEL SYNDROME"

*Submitted in partial fulfilment of
requirements for the degree of*

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of

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CERTIFICATE

This is to certify that this dissertation entitled “**EVALUATION OF CLINICAL, ELECTROPHYSIOLOGICAL, AND RADIOLOGICAL PROFILE OF CARPAL TUNNEL SYNDROME**” submitted by **Dr. K. GANESAN** appearing for **D.M. Neurology** Degree (Branch - I) examination in **August 2009** is a bonafide record of work done by him under my direct guidance and supervision in partial fulfillment of regulations of the Tamil Nadu Dr. M.G.R. Medical University, Chennai. I forward this to the Tamil Nadu Dr.M.G.R. Medical University, Chennai, Tamil Nadu, India.

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DECLARATION

I solemnly declare that the dissertation titled “**EVALUATION OF CLINICAL, ELECTROPHYSIOLOGICAL, AND RADIOLOGICAL PROFILE OF CARPAL TUNNEL SYNDROME**” is done by me at Institute of Neurology, Madras Medical College & Govt. General Hospital, Chennai, during 2007-2009 under the guidance and supervision of **Prof. V. NATARAJAN, M.D., D.M.**,

The dissertation is submitted to The Tamil Nadu Dr. M.G.R. Medical University towards the partial fulfillment of requirements for the award of **D.M., degree in Neurology.**

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INTRODUCTION

Carpal tunnel syndrome (CTS) is the most commonly-encountered entrapment neuropathy with an incidence of 139 per 100,000 person-years for men and 506 per 100,000 person-years for women ¹²⁵. The symptoms and signs are caused by compression of the median nerve along the carpal tunnel, which is formed on the distal, medial, and lateral sides by the carpal bones and on the volar surface by the deep transverse carpal ligaments.

The classic symptoms of CTS are numbness and paraesthesia in the first three fingers of the hand, which is commonly exacerbated at night ¹²⁵. The diagnostic signs include sensory loss along the lateral aspect of the hand, motor weakness and wasting of abductor pollicis brevis (APB) muscle, and eliciting Tinel's and Phalen's sign at the wrist.

The nerve conduction study (NCS) study is a definite diagnostic test for CTS with high degree of sensitivity and specificity. This test demonstrates a distal lesion of the median nerve and excludes other peripheral conditions resulting in similar symptoms ^{125,126}. Ultrasonography is also an upcoming tool in the diagnosis of carpal tunnel syndrome with equal sensitivity as electrophysiological studies.

The present study is being done to evaluate the clinical profile, electrophysiological severity and radiological profile in patients with symptoms and signs suggestive of carpal tunnel syndrome.

AIM OF THE STUDY

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- ❖ To correlate the clinical, electrophysiological and radiological findings in carpal tunnel syndrome.
- ❖ To study the etiologic profiles in adult patients aged more than 12 years of age with carpal tunnel syndrome.
- ❖ To detect the sensitivity and specificity of various neurophysiological tests and ultrasonogram of wrists in patients with carpal tunnel syndrome.
- ❖ To identify the frequency of subclinical impairment on the unaffected side and identify other pathologies which could cause symptoms of carpal tunnel syndrome.

MATERIALS AND METHODS

MATERIALS AND METHODS

- Place of study** : **Institute of Neurology,**
Government General Hospital,
Madras Medical College,
Chennai -3.
- Type of study** : Cross sectional, Clinical and Investigatory Study
- Duration of study** : 2 years (March 2007 to February 2009).
- Ethical committee** : Present dissertation was approved
by the Institutional Ethics Committee.
- Consent** : Informed written consent was
obtained from all the participants.
- Collaborating
department** : **Barnard Institute of Radiology**
Government General Hospital,
Madras Medical College,
Chennai -3.

Case selection

Patients with clinical symptoms suggestive of carpal tunnel syndrome of both sexes were selected from the Neurology OPD, Institute of Neurology, Madras Medical College, Chennai- 3.

A. Inclusion criteria

1. Patients with clinical symptoms and signs of carpal tunnel syndrome of both sexes aged above 12 years.

2. Patients with polyneuropathies associated with clinical symptoms and signs of carpal tunnel syndrome of both sexes aged above 12 years.

B. Exclusion criteria:

1. Patients with age less than 12 years.
2. Patients with incomplete clinical profiles / diagnostic records were excluded.

Limitation of the study:

- Due to technical and administrative reasons, interobserver bias in the ultrasound examination of wrists could not be excluded.

Materials:

The total number of 96 patients who satisfied the inclusion criteria formed the final materials of the present study.

Methods:

All subjects gave their informed consent prior to the study. History taking and detailed neurological evaluation was done in all patients, followed by blood investigations, electrophysiology, and ultrasonogram of both wrists. The findings were entered in the proforma. (Copy of proforma enclosed)

Blood Investigations done included:

1. GTT
2. Thyroid function tests
3. Serum rheumatoid factor

Electrophysiology:

The nerve conduction studies (NCS) and electromyography (EMG) were selected

- 1) To demonstrate a distal lesion of the median nerve
- 2) To exclude other peripheral conditions that can result in similar symptoms like proximal median neuropathy, C6-C7 radiculopathy, lesions of the brachial plexus and an attempt was made to correlate the sensitivity and specificity of various neurophysiological tests

Nerve Conduction Study Protocol

A. Routine studies

- 1) Median motor study recording abductor pollicis brevis. Stimulating wrist and antecubital fossa
- 2) Ulnar motor study recording abductor digiti minimi. Stimulating wrist, below groove, and above groove

- 3) Median and ulnar F responses
- 4) Median sensory response, recording digit 2 or 3, stimulating wrist
- 5) Ulnar sensory response, recording digit 5, stimulating wrist

B. Median-versus-ulnar comparisons

- 1) Comparison of the median lumbrical and ulnar interossei distal motor latencies, stimulating the median and ulnar wrist one at a time at identical distances (8-10 cm), recording with the same electrode over the 2L/interossei
- 2) Comparison of the median and ulnar digit 4 sensory latencies, stimulating the median and ulnar wrist one at a time at identical distances (11-13 cm) and recording digit 4
- 3) F wave latencies of median nerve on symptomatic side were compared with ulnar F wave latencies of that side and with the median nerve F latencies on the asymptomatic side.

If motor & sensory nerve conduction studies showed reduced amplitude of distal CMAP's and SNAP's of median nerve, Palm vs wrist comparison studies were done to differentiate between axonal and conduction block.

C. Palm wrist comparison studies

- 1) Palm vs wrist motor conduction of median nerve looking for significant increase in CMAP amplitude (distal/proximal ratio >1.2)

2) Palm vs wrist sensory conduction of median nerve looking for significant increase in SNAP amplitude (distal/proximal ratio >1.6)

D. If the ulnar motor and/or sensory studies also are abnormal, **additional nerve** (motor, sensory, F wave) **conduction studies** were done in both upper limbs and lower limbs to rule out brachial plexopathy, C6-C7 radiculopathy, and polyneuropathies.

E. Terminal latency index (TLI):

TLI was calculated in all patients using the following formula

$$\text{TLI} = \frac{\text{Terminal distance (mm)}}{\text{CV (m/sec)} \times \text{TL (msec)}}$$

The sensitivity and specificity of various neurophysiological tests were studied.

Additional EMG studies were done in cases of axonal changes on median motor stimulation and in cases of suspected cervical radiculopathies.

Electromyography protocol

Electromyography of following muscle groups was done.

- 1) Abductor pollicis brevis (APB)
- 2) C6-C7 muscles (e.g., pronator teres, triceps brachii, extensor digitorum communis) to exclude a cervical radiculopathy

If APB is abnormal, the following additional muscles were sampled:

- 1) One proximal median muscle (e.g., flexor carpi radialis, pronator teres, flexor pollicis longus) to exclude a proximal median neuropathy
- 2) Two other non median, lower trunk/C8-T1 muscles (e.g., first dorsal interosseous, extensor indicis proprius) to exclude a lower trunk brachial plexopathy, polyneuropathy, or C8-T1 radiculopathy

Recorders and Medicare systems electrophysiology machine was used. Conventional methods for the measurement of nerve conduction were employed. The room temperature was kept between 22-24⁰ C, and before the testing it was made sure that the limbs were warm enough; at least 32⁰ C.

NERVE CONDUCTION STUDIES – NORMAL ADULT VALUES.

1. UPPER EXTREMITY – MOTOR

Nerve	Recording site	Distal latency (ms)	Amplitude (mv)	Conduction velocity (m/s)	Distal distance (cm)	F response latency(ms)
Median	APB	≤ 4.4	≥ 4.0	≥ 49	7	≤ 31
Ulnar	ADM	≤ 3.3	≥ 6.0	≥ 49	7	≤ 32

APB –abductor pollicis brevis, ADM- abductor digiti minimi

2. UPPER EXTREMITY - ANTIDROMIC SENSORY

Nerve	Recording site	Distal latency (ms)	Amplitude (μv)	Conduction velocity (m/s)	Distal distance (cm)
Median	Digit 2	≤ 3.5	≥ 20	≥ 50	13
Ulnar	Digit 5	≤ 3.1	≥ 17	≥ 50	11

3. MEDIAN – ULNAR INTERNAL COMPARISON STUDIES

Study*		Significant Latency Difference (ms)#
Median motor vs Ulnar motor	Wrist to 2 nd lumbrical vs Wrist to interossei	≥ 0.5
Median sensory vs Ulnar sensory	Wrist to digit 4 vs Wrist to digit 4	≥ 0.5

* - For each paired study, identical distances are used for both the median and ulnar study (8-10 for motor & 11-13 for sensory).

- Value that exceeds these cutoffs implies focal slowing and is useful in electrodiagnosis of median neuropathy across the carpal tunnel.

4. MEDIAN PALMAR AND WRIST STIMULATION COMPARISON STUDIES

Study*		Significant Palm to wrist Amplitude Ratio#
Median motor	Wrist to APB	>1.2
Median motor	Palm to APB	
Median sensory	Wrist to digit 2	>1.6
Median sensory	Palm to digit 2	

*- Both studies are supramaximal stimulation

- Values that exceed these cutoffs imply some element of conduction block of the median nerve across the carpal tunnel. Values below these cutoffs on distal stimulation imply secondary axonal loss.

5. LOWER EXTREMITY - MOTOR

Nerve	Recording site	Distal latency (ms)	Amplitude (mv)	Conduction velocity (m/s)	F response latency(ms)
Peroneal	EDB	≤ 6.5	≥ 2.0	≥ 44	≤ 56
Tibial	AHB	≤ 5.8	≥ 3.0	≥ 41	≤ 56

EDB- extensor digitalis brevis, AHB – abductor hallucis brevis

6. LOWER EXTREMITY - ANTIDROMIC SENSORY

Nerve	Recording site	Distal latency (ms)	Amplitude (μv)	Conduction velocity (m/s)
Sural	Posterior Ankle	≤ 4.4	≥ 6	≥ 40

Electrophysiological Parameters Taken To Confirm CTS Were

Median studies:

- Prolonged distal motor latency > 4.4 ms
- Prolonged distal sensory (digit2) latency > 3.5 ms
- Prolonged minimum F wave latency > 31 ms or F waves latency difference between ipsilateral median and ulnar nerves > 2 ms or F waves latency difference between median nerves of symptomatic side and contralateral side > 2 ms.
- Terminal latency index < 0.33

Comparison studies:

- Difference between distal motor latency of median and ulnar nerves > 1.1 ms
- Difference between distal sensory latency of median and ulnar nerves > 0.2 ms
- Difference between median and ulnar digit 4 sensory latencies ≥ 0.5 ms

- Difference between median lumbrical and ulnar interossei distal motor latencies \geq 0.5ms

Palm and wrist stimulation comparison

In case of decreased distal amplitude of CMAP'S (<4.2mv) or SNAP's (<20 μ v) of median nerve stimulation at wrist, palm and wrist stimulation comparison studies done.

Abnormal if

- Significant increase in amplitude of CMAP's on distal palm stimulation ie, if distal/proximal ratio is >1.2(segmental demyelination with conduction block) or if there is no increase in ratio >1.2 (secondary axonal loss).
- Significant increase in amplitude of SNAP's on distal palm stimulation ie, if distal/proximal ratio is >1.6(segmental demyelination) or if there is no increase in ratio >1.6 (secondary axonal loss).

The Severity of Neurophysiologic Impairment in CTS.

The severity of neurophysiologic CTS impairment is assessed by the neuro physiologic classification by Padua et al¹²⁷.

CTS hands are divided into six groups on the basis of the neurophysiologic findings.

Neurophysiological classification (Padua L et al) ¹²⁷,

STAGING	SENSORY NCS	MOTOR NCS
NEGATIVE	Normal	Normal
MINIMAL	Abnormal segmental and /or comparative test only	Normal distal motor latency
MILD	Abnormal digit-wrist conduction	Normal distal motor latency
MODERATE	Abnormal digit-wrist conduction	Abnormal distal motor latency
SEVERE	Absence of response	Abnormal distal motor latency
EXTREME	Absence of response	Absence of response

In the patients with bilateral CTS, the neurophysiological grade in the more severely affected hand was noted.

ULTRASONOGRAM WRISTS:

Ultrasonography protocol:

All patients underwent high-resolution real-time sonography of the carpal tunnel (both hands) using 7.5 MHz linear array transducer. In initial part of the study, ALOKA ultrasonography machine and in later part of the study SIEMANS ultrasonography machine was used.

The sonographic examination was performed with the patient seated in a comfortable position facing the sonographer, with the forearm resting on the table and the palm

facing up in the neutral position. The volar wrist crease was used as an initial external reference point, with subsequent modifications during

scanning, using carpal bony landmarks and internal reference points. The full course of the median nerve in the carpal tunnel was assessed in both transverse and longitudinal planes. The median nerve is located superficial to the echogenic flexor tendons and its size, shape, echogenicity, and relationship to the surrounding structures and overlying retinaculum were noted. The amount of synovial fluid and the presence or absence of masses was noted.

The continuity of the median nerve and any area of constriction were assessed in both the longitudinal and transverse planes. Measurements were taken for the median nerve at the carpal tunnel inlet proximally.

The mean cross-sectional area of the median nerve was measured by tracing with electronic calipers around the margin of the nerve at the time of sonography (direct tracing). The flattening ratio (defined as

the ratio of the major axis of the median nerve to its minor axis) was also assessed at the tunnel inlet.

Carpal tunnel syndrome was diagnosed when there was

- (1) Increased cross-sectional area of the median nerve at the pisiform bone and/or at the hamate bone $> 0.09 \text{ cm}^2$.
- (2) Increased flattening ratio of the median nerve at the hamate bone > 2 , or
- (3) Increased palmar displacement of the flexor retinaculum $> 1 \text{ mm}$.

Observation and results

Observation and results

A total of 96 patients aged above 12 years who came to Neurology OPD, Government General Hospital, Chennai between March 2007 to February 2009 with clinical features suggestive carpal tunnel syndrome and who satisfied the inclusion and exclusion criteria were included in this study.

- ❖ Total number of patients - 96
- ❖ Male patients - 28(29.2%)
- ❖ Female patients - 68(70.8%)
- ❖ Male: Female ratio - 1:2.42

AGE DISTRIBUTION:

- The maximum numbers of patients were in the age group between 40 and 49 years, followed by the age group between 50 and 59, and 30 and 39 years.

Table 1 shows the age distribution in this study

TABLE - 1: AGE DISTRIBUTION

<i>Age group in years</i>	<i>No. of Patients</i>	<i>% of Total Patients (96)</i>
13-19	-	-
20-29	10	10.41
30-39	19	19.80
40-49	32	33.33
50-59	24	25.00
60-69	7	7.29
70-79	3	3.13
80 & above	1	1.04
Total	96	100

SEX Distribution:

- There were 28 males (29.2%) and 68 females (70.8%) among the 96 patients in this study.

TABLE 2: SEX DISTRIBUTION IN THIS STUDY

<i>Sex</i>	<i>No. of Patients</i>	<i>% of Total Patients (490)</i>
Males	28	29.2
Females	68	70.8
Total	96	100

Age and Sex distribution:

- Males and females predominated in the age group between 40 and 49 years.
- Around two thirds of males (64.3%) were in the age group between 30 and 59 years and two-third of females (63.2%) were in the age group between 40 and 59 years. The Table 3 shows age distribution based on sex.

TABLE 3: AGE DISTRIBUTION BASED ON SEX

<i>Age group in years</i>	<i>Males (%)</i>	<i>Females (%)</i>
13-19	-	-
20-29	3(10.7)	7(10.3)
30-39	5(17.9)	14(20.6)
40-49	8(28.6)	24(35.3)
50-59	5(17.9)	19(27.9)
60-69	4(14.3)	3(4.4)
70-79	2(7.0)	19(1.5)
80 & above	1(3.6)	-
Total	28(100)	68(100)

DURATION OF SYMPTOMS

- Most of the males (35.7%) and females (51.5%) had symptom duration of less than 6 months.

TABLE 4: DURATION OF SYMPTOMS

Duration	Males (%)	Females (%)	Total (%)
< 6 months	10(35.7)	35(51.5)	45(46.9)
6 months – 1 year	9(32.1)	23(33.8)	32(33.3)
1 year – 2 years	4(14.3)	6(8.8)	10(10.4)
2 years and above	5(17.9)	4(5.9)	9(9.4)
Total	28	68	96

DURATION OF SYMPTOMS WITH CTS SEVERITY

- Most of the patients with more than 2 years of symptoms had extreme CTS (88%). Those patients with duration less than 6 months had minimal to mild CTS. As the duration increases, there was a progression from mild to extreme CTS.

TABLE: 5 DURATION OF SYMPTOMS WITH CTS SEVERITY

NCS severity	< 6 months			6 mths – 1 year			1 year - 2years			2 yrs. & above			TOTAL		
	M	F	T	M	F	T	M	F	T	M	F	T	M	F	T
1	5	15	20	-	-	-	-	-	-	-	-	-	5	15	20
2	5	16	21	-	1	1	-	-	-	-	-	-	5	17	22
3	-	4	4	8	22	32	-	-	-	-	-	-	8	26	34
4	-	-	-	1	-	1	4	6	10	1	-	1	6	6	12
5	-	-	-	-	-	-	-	-	-	4	4	8	4	4	8
Total	10	35	45	9	23	32	4	6	10	5	4	9	28	68	96

PARESTHESIAS

- Paresthesias were seen in more than two third (70.8%) of patients. 78.6%

of males and 67.6 % of females with CTS had paresthesias. In patients with paresthesias, median distribution of paresthesias was seen in majority of the patients, 54.5% of males, and 52.1% of females.

TABLE 6: DISTRIBUTION OF PARESTHESIAS

Distribution	Male (%)	Female (%)	Total (%)
Median	12(54.5%)	24 (52.1%)	36(52.9%)
Extra median	10(45.5%)	22(47.8%)	32(47.1%)
Total	22/28(78.6%)	46/68(67.6%)	68/96(70.8%)

- In patients with paresthesias more than two thirds of them had nocturnal paresthesias, 63.6% of males, and 60.9% of females.

TABLE 7: OCCURRENCE OF PARESTHESIAS

Occurrence of paresthesias	Male	Female	Total
Nocturnal	14(63.6%)	28(60.9%)	42(61.8%)
Nocturnal +diurnal	8(36.4%)	18(39.1%)	26(38.2%)
Total	22	46	68

NUMBNESS

- Numbness in median distribution was seen in one third (33.3%) of patients.

TABLE 8: NUMBNESS

Distribution	Male	Female	Total
Median	7(25%)	25(36.8%)	32(33.3%)

ARTHRALGIA /JOINT SWELLING

- Nearly one sixth (15.6%) of patients had joint pains and all were females.

TABLE 9: JOINT PAIN AND SWELLING

	Male	Female	Total
Joint pains & swelling	--	15	15(15.6%)

OCCUPATION

- Majority of females (67.6%) having symptoms were involved in household works. Nearly half of males (60.7%) with symptoms were doing manual work

TABLE 10: OCCUPATION WISE DISTRIBUTION

Occupation	Male	Female	Total
House wife	-	46(67.6%)	46(47.9%)
Construction labour	8(28.6%)	8(11.8%)	16(16.7%)
Agricultural labour	5(17.9%)	-	5(5.2%)
Mechanic	4(14.2%)	-	4(4.2%)
Teacher	2(7.1%)	4(5.9%)	6(6.3%)
College student	1(3.6%)	3(4.4%)	4(4.2%)
Computer operator	1(3.6%)	4(5.9%)	5(5.1%)
Staff nurse	-	3(4.4%)	3(3.1%)
Others	7(25%)	-	7(7.3%)
Total	28	68	96

SENSORY LOSS

- Nearly one third (40.6%) of patients had median distribution of sensory

loss. Two patients had C6-C7 root distribution of sensory loss while ten patients had glove and stocking sensory loss.

TABLE 11: SENSORY LOSS

Distribution	Male	Female	Total
Median	9(32.1%)	30(44.1%)	39(40.6%)
C6-C7 root	1(3.6%)	1(1.5%)	2(2.1%)
Glove & stocking	7(25.0%)	3(4.4%)	10(10.4%)

WEAKNESS OF MEDIAN INNERVATED HAND MUSCLES

- Weakness of median innervated hand muscles (1&2 lumbricals, opponens pollicis, abductor pollicis brevis, flexor pollicis brevis) was seen in one third (39.3%) of males and one fourth (25%) of females.

TABLE 12: WEAKNESS OF MEDIAN INNERVATED HAND MUSCLES

Male	Female	Total
11(39.3%)	17(25%)	28(29.2%)

WASTING OF THENAR MUSCLES

- Wasting of thenar muscles was seen in 10.7% of males and 5.9% of females.

TABLE 13: WASTING OF THENAR MUSCLES

Male	Female	Total
3(10.7%)	4(5.9%)	7(7.3%)

TINEL'S SIGN

- Tinel's sign was positive in 38.5% of individuals. Tinel's sign was positive in nearly one third of males (32.1%) and females (29.2%).

TABLE 14: TINEL'S SIGN

Tinel's sign	Male	Female	Total
Positive	9(32.1%)	28(29.2%)	37(38.5%)

PHALEN'S TEST

- Phalen's test was positive in 64.5% of individuals. Phalen's test was positive in nearly two third of males (64.3%) and females (64.7%).

TABLE 15: PHALEN'S TEST

Phalen's test	Male	Female	Total
Positive	18(64.3%)	44(64.7%)	62(64.5%)

BLOOD INVESTIGATIONS

- Glucose tolerance test was done in all patients and 13(13.5%) patients were diagnosed as diabetics (whose fasting blood glucose >126mg% and/or 2 hours post meal blood glucose > 200mg %).
- Diabetes was most commonly seen in males (28.6%) than females (7.4%).

- Thyroid function tests were done in all patients and 7(7.3%) patients were found to have hypothyroidism.
- Rheumatoid factor estimation was done in all patients and it was exclusively positive in 18.8 %.(all were females).

TABLE 16: BLOOD INVESTIGATIONS

Investigation	Male	Female	Total
Diabetes	8(28.6%)	5(7.4%)	13(13.5%)
Hypothyroidism	1(3.6%)	6(8.8%)	7(7.3%)
RA factor + ve	-	18(26.5%)	18(18.8%)

SYMPTOMS

- Nearly half of the patients had bilateral symptoms on presentation.

TABLE 17: SYMPTOMS

Symptoms	Male	Female	Total
Bilateral	14(50%)	38(55.9%)	52(54.2%)
Unilateral	14(50%)	30(44.1%)	44(45.8%)
Total	28	68	96

UNILATERAL SYMPTOMS

- Nearly two thirds (65.9%) of patients had symptoms in dominant hand and one third (34.1%) of patients in their nondominant hands

TABLE 18: UNILATERAL SYMPTOMS

Symptoms	Male	Female	Total
Dominant	10	19	29(65.9%)
Non dominant	4	11	15(34.1%)
Total	14	30	44

ETIOLOGY

- Majority of patients had no obvious cause (44.8%) for CTS and were considered to be idiopathic.
- Diabetes was the 2nd common cause in males (28.6%) whereas rheumatoid arthritis was the 2nd common cause among females (26.5%).
- Polyneuropathy was the third common cause of CTS in males (14.3%) and hypothyroidism in females (8.8%).
- C6-C7 radiculopathy was mimicking CTS in one male and one female patient. C8 -T1 radiculopathy was seen in another male patient.
- Previous history of trauma with injury to the wrist was obtained in three male patients.
- Out of 3 pregnant females 1 was a primigravida with pregnancy induced hypertension (PIH) and other 2 were multigravida without any comorbid illness.

TABLE 19: ETIOLOGY

Etiology	Male	Female	Total
Idiopathic	10(35.7%)	33(48.5%)	43(44.8%)
Diabetes	8(28.6%)	5(7.4%)	13(13.5%)
Rheumatoid arthritis	-	18(26.5%)	18(18.8%)
Hypothyroidism	1(3.6%)	6(8.8%)	7(7.3%)
Pregnancy	-	3(4.4%)	3(3.1%)
Polyneuropathy	4(14.3%)	2(2.9%)	6(6.3%)
Trauma	3(10.7%)	-	3(3.1%)
Radiculopathy	2(7.1%)	1(1.5%)	3(3.1%)
Total	28	68	96

BILATERAL CTS WITH ASYMPTOMATIC INVOLVEMENT OF OTHER HAND DETECTED ON NERVE CONDUCTION STUDIES

- 19 out of 41(46.3%) patients with unilateral symptoms showed subclinical CTS in the other hand also after nerve conduction studies. After NCS nearly three fourths (76.3%) of patients had bilateral carpal tunnel syndrome.
- Almost two thirds of male (69.2%) and three fourths of female (79.1%) patients had bilateral carpal tunnel syndrome (CTS) after electrophysiology. 3 patients with radiculopathy mimicking CTS with unilateral symptoms were excluded.

TABLE 20:

BILATERAL CTS AFTER NERVE CONDUCTION STUDIES

Symptoms	Male	Female	Total
Bilateral CTS	14(50.0%))	38(55.8%))	52(54.2%)
Asymptomatic CTS in other hand detected by NCS in patients with unilateral symptoms	4	15	19
Total	18(69.2%))	53(79.1%))	71(76.3%)

UNILATERAL CTS AFTER NERVE CONDUCTION STUDIES

Unilateral CTS	8	14	22(23.7%)
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- So 71 patients (18 males and 53 females) had bilateral CTS and 22 patients (8 males and 14 females) had unilateral CTS.
- So a total of 186 hands were examined and 164 confirmed hands with CTS were included in the study.

ELECTROPHYSIOLOGICAL GRADING OF SEVERITY OF CARPAL TUNNEL SYNDROME (Padua et al¹²⁷)

- In the patients with bilateral CTS, the electrophysiological grade in the more severely affected hand was noted.

TABLE 21: ELECTROPHYSIOLOGICAL GRADING OF SEVERITY OF CTS

Staging	Male	Female	Total
Minimal	5	15	20 (21.5%)
Mild	5	17	22(23.7%)
Moderate	7	25	32(34.4%)
Severe	6	6	12(12.9%)
Extreme	3	4	7(7.5%)
Total	26	67	93

A. IDIOPATHIC

- Nearly two third (62.7%) of the idiopathic carpal tunnel syndrome patients had mild to moderate CTS

TABLE: 22. GRADING IN IDIOPATHIC PATIENTS

Staging	Male	Female	Total
Minimal	3(30%)	7(21.2%)	10(23.3%)
Mild	2(20%)	11(33.3%)	13(30.2%)
Moderate	2(20%)	12(36.4%)	14(32.5%)
Severe	2(20%)	2(6.1%)	4(9.3%)
Extreme	1(20%)	1(3.0%)	2(4.7%)
Total	10	33	43

B.RHEUMATOID ARTHRITIS

- Nearly two third (66.6%) of CTS patients with rheumatoid arthritis had mild to moderate CTS

TABLE: 23. GRADING IN RHEUMATOID ARTHRITIS PATIENTS

Staging	Male	Female	Total
Minimal	-	4	4(22.2%)
Mild	-	4	4(22.2%)
Moderate	-	8	8(44.4%)
Severe	-	1	1(5.6%)
Extreme	-	1	1(5.6%)
Total	-	18	18

C.DIABETES

- Nearly one third (31%) of patients with diabetes had severe to extreme CTS.

TABLE: 24: GRADING IN DIABETES PATIENTS

Staging	Male	Female	Total
Minimal	2	1	3(23.0%)
Mild	2	1	3(23.0%)
Moderate	2	1	3(23.0%)
Severe	1	1	2(15.5%)
Extreme	1	1	2(15.5%)
Total	8	5	13

D.HYPOTHYROIDISM

- 6 out of the total 7 patients with hypothyroidism were females.

TABLE 25: GRADING IN HYPOTHYROID PATIENTS

Staging	Male	Female	Total
Minimal	-	2	2(28.7%)
Mild	-	1	1(14.2%)
Moderate	1	1	2(28.7%)
Severe	-	1	1(14.2%)
Extreme	-	1	1(14.2%)
Total	1	6	7

E.POLYNEUROPATHY

- More than 80% patients with polyneuropathy have moderate to severe CTS.

TABLE 26: GRADING IN POLYNEUROPATHY PATIENTS

Staging	Male	Female	Total
Minimal	-	-	-
Mild	1	-	1(16.7%)
Moderate	2	1	3(50.0%)
Severe	1	1	2(33.3%)
Extreme	-	-	-
Total	4	2	6

F.PREGNANCY

- Of 3 pregnant women with CTS 1 had bilateral minimal CTS and 2 had bilateral moderate CTS.

TABLE: 27: GRADING IN PREGNANT PATIENTS

Staging	Female	Total
Minimal	1	1
Mild	-	-
Moderate	2	2
Severe	-	-
Extreme	-	-
Total	3	3

G. TRAUMA

- Of 3 patients with trauma 1 had healed Colles' fracture right wrist and 2 had history of trauma wrists in their symptomatic hands.

TABLE: 28 TRAUMA

Staging	Male	Female	Total
Minimal	-	-	-
Mild	-	-	-
Moderate	-	-	-
Severe	2	-	2
Extreme	1	-	1
Total	3	-	3

SENSITIVITY OF VARIOUS ELECTROPHYSIOLOGICAL STUDIES

- Of the various electrophysiological tests done, digit 4 sensory median ulnar comparison study was most sensitive (92.1%) followed by F wave comparison studies (87.2%).terminal latency index was least sensitive(54.8%).

TABLE: 29 A. DIGIT 2 SENSORY NCS

SNAP's	
Prolonged distal latency/ Decreased amplitudes	112
Absent SNAP's	19
Total	131(79.9%)

TABLE: 29 B. MEDIAN MOTOR NCS

CMAP's	
Prolonged distal latency/ Decreased amplitudes	92
Absent CMAP's	7
Total	99(60.3%)

TABLE: 29 C. F WAVES

F WAVE	
Abnormal F-wave latency	136
Absent CMAP's	7
Total	143(87.2%)

TABLE: 29 D. DIGIT 4 SENSORY

SNAP	
Abnormal (>0.5msec difference)	132
Absent SNAP's	19
Total	143(92.1%)

TABLE: 29 E. 2nd LUMBRICAL/INTEROSSEI

Abnormal (>0.5msec difference)	106(64.6%)
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TABLE: 29 F. TERMINAL LATENCY INDEX

Abnormal	90(54.8%)
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EMG

- 3 out of 7 patients with extreme CTS showed neurogenic pattern on EMG of Abductor pollicis brevis.
- Of the 3 patients 1 was a male patient with trauma, another was a male patient with no obvious cause and the third was a female with rheumatoid arthritis.
- The remaining 4 patients (1 male and 3 female) patients had conduction block which was confirmed by palm and wrist stimulation motor comparison study which showed distal/proximal ratio difference in amplitude of >1.2.

- 2 patients (1 male & 1 female) showed neurogenic pattern on EMG of the following muscles (Pronator teres & triceps) and were found to have C6-C7 radiculopathy.
- 1 male patient showed neurogenic pattern in 1st dorsal interossei, abductor digiti minimi in addition to abductor pollicis brevis and was found to have C8-T1 radiculopathy.

ULTRASONOGRAM BOTH WRISTS

- Ultrasonogram showed evidence of CTS in 59 of 93(63.5%) patients with CTS.
- In addition to increased median nerve cross sectional area ,USG also showed swelling of tendon sheaths in 12 female patients with rheumatoid arthritis.

TABLE: 30. ULTRASONOGRAM

USG	Male	Female	Total
Abnormal (Median nerve Cross sectional area >0.09 cm ² at carpal tunnel inlet)	17	42	59 (63.5%)
Normal	9	25	34
Total	26	67	93

Electrophysiological staging wise USG both wrists

- Ultrasonography detected abnormalities in late stages of CTS better than early stages

TABLE 31: Electrophysiological staging wise USG both wrists

Staging	Abnormal			Normal		
	M	F	T	M	F	T
1	1	1	2	4	14	18
2	5	15	20	-	2	2
3	4	16	20	3	9	12
4	4	6	10	2	-	2
5	3	4	7	-	-	-
Total	17	42	59	9	25	34

DISCUSSION

DISCUSSION

SEX DISTRIBUTION

The present study, showed a female preponderance in carpal tunnel syndrome patients. Of 96 patients, 70.8 were females and 29.2% were males. The male: female ratio was 1: 2.42.

According to the study by Mondelli et al, carpal tunnel syndrome (CTS) is the most commonly-encountered entrapment neuropathy with an incidence more in women than men.⁹⁶ In the studies done by Phalen et al, and Radecki P et al.^{97,98} 3-4 times higher incidence of carpal tunnel syndrome in females have been quoted. In a recent surveillance study from Canterbury and Huddersfield, UK, Bland *et al*⁹⁹ reported female to male ratio of 2.07. The present study showed almost similar findings as the above studies.

AGE DISTRIBUTION

Maximum number of patients in our study were in the age group between 40 and 49 years (33.33%), followed by the age group between 50 and 59 years (25.00%).

According to the study by Tay L B¹⁰⁰ et al, more than two-thirds of the patients in their study were between 40 and 70 years of age.

In the present study also more than two thirds (65.6%) belong to 40 – 70 years of age.

DURATION OF SYMPTOMS

In the present study, 46.9% of individuals had symptom duration of less than 6 months. These patients with duration of less than 6 months had minimal to mild CTS. Most of the patients with more than 2 years of symptoms had extreme CTS (88%). As the duration increases there was a progression from mild to extreme CTS.

In the study by Urkude et al¹⁰⁰ the median duration of symptoms for mild, moderate, and severe CTS were two months, four months and 12 months, respectively, suggesting the increasing severity of CTS with longer duration of symptoms.

The above finding very well correlates with the present study that severity of CTS increases with the duration of symptoms.

PARESTHESIAS

Paresthesias were seen in more than two third (70.8%) of patients. 78.6% of males and 67.6 % of females with CTS had paresthesias. In patients with paresthesias, median distribution of paresthesias was seen in majority (52.9%). More than two thirds (61.8%) of patients with paresthesias had nocturnal paresthesias. ie, 63.6% of males, and 60.9% of females.

In Kendall's¹⁰³ series of 327 patients, 313 (95.7%) reported paresthesia; 118 (38%) reported nocturnal symptoms only, 178 (58%) reported symptoms during the day and night, but worse at night.

Symptoms of nocturnal paraesthesia are reported to be 51-96% sensitive and 27-68%

specific as per the studies by Szabo RM et al, Buch-Jaeger et al, and Gupta SK et al.⁵⁸⁻⁶¹.

In another study by **Tay L B**¹⁰⁰et al. paresthesia was present in 70.1% patients.

The present study showed similar results as the study by Tay L B et al.

NUMBNESS

Numbness in median distribution was seen in one third (33.3%) of patients.

Numbness was present in 19.4% patients in a study by **Tay L B**¹⁰⁰et al.

OCCUPATION

Majority of females (67.6%) having symptoms were housewives engaged in household works. Nearly half of males (60.7%) with symptoms were doing manual work

Experimental studies have shown a higher incidence of CTS in workers who are involved in high force and repetitive work compared to workers who are not.²⁸⁻³¹.

Silverstein BA et al²⁸ noted a prevalence of 5.6% among workers in high force and high repetitive jobs compared to 0.6% among workers in low force and low repetitive jobs.

Palmer *et al*⁵³, found that the regular use of hand-held vibrating tools increased the risk of CTS by more than 2-fold. They also found substantial evidence for high risk of CTS in occupations requiring high repetitive flexion and extension at wrist and also forceful grip⁵³. However, they did not find evidence

between the work on keyboard and computers and CTS.

Similar to the above studies, our study also showed majority of the patients with symptoms of CTS were house wives and manual laborers who were involved in high force and repetitive jobs. Our studies also showed that, relatively few (5.1%) of the total patients were computer operators.

SENSORY LOSS, WEAKNESS, AND WASTING

In our study, nearly one third (40.6%) of patients had median distribution of sensory loss. Weakness of median innervated hand muscles (1&2 lumbricals, opponens pollicis, abductor pollicis brevis, flexor pollicis brevis) was seen in one third (29.2%) of patients. Wasting of thenar muscles was seen in 7.3% of patients.

Thenar atrophy is a late sign and signifies significant functional loss. Finger weakness associated with an inability to pinch or frequent dropping of grasped objects, follows involvement of the motor component. Long-term involvement leads to thenar muscle atrophy with associated loss of thumb abduction and opposition strength. Thenar atrophy is seldom noticed by patients and may not be obvious even to the examiner when examined by looking down onto the palm. However, it will be readily appreciated by comparing both palms together.

Tay L B¹⁰⁰ et al. in their study found APB weakness in 10.5% patients and APB muscle wasting in 10.4% patients.

In Phalen's ¹⁵series the atrophy of abductor pollicis brevis, opponens pollicis, and flexor pollicis brevis was noted in 41% of hands.

In our study, patients showed more of weakness than wasting of thenar muscles.

TINEL'S SIGN & PHALEN'S TEST

In the present study, Tinel's sign was positive in 38.5% of individuals. Phalen's test was positive in 64.5% of individuals.

The Tinel's sign is associated with sensitivities of 23% to 67%, and specificities of 55% to 100%^{59,60,65,67-71} according to various studies by Kuhlman KA et al, Buch-Jaeger et al, and Katz JN. et al.

In a review, Kushner *et al*⁶⁷ summarized the frequency of Tinel's sign and reported that it is positive from 8% to 100% of CTS patients. Tinel's sign is the least accurate test according to Mondelli *et al*⁷¹, who did not find a combination of signs more useful than a single sign alone.

The reported sensitivity of Phalen's test ranges between 10% and 91% and specificity between 33% and 100%^{59, 60, 68, 69, 73-75}.

Tinel's and Phalen's sign were positive in 33.6% and 23.9% of patients respectively in the study by Tay L B¹⁰⁰ et al.

In our study sensitivity of the Tinel's sign and Phalen's test are 38.5% and 64.5% respectively.

BILATERAL VS UNILATERAL CTS

In our study, nearly half (54.2%) of the patients had bilateral symptoms on

presentation. 50% of male patients and 55.9% of female patients had bilateral symptoms at presentation. Nearly two thirds (65.9%) of patients had symptoms in dominant hand and one third (34.1%) of patients in their nondominant hands.

Based on NCS nearly three fourths (76.3%) of patients had bilateral carpal tunnel syndrome and one fourth(23.7%) of patients had unilateral CTS. 19 out of 41 (46.3%) patients with unilateral symptoms showed subclinical CTS in the other hand also on nerve conduction studies.

After electrophysiology nearly two thirds of male (69.2%) and three fourths of female (79.1%) patients were detected to have bilateral carpal tunnel syndrome (CTS). 3 patients with radiculopathy mimicking CTS with unilateral symptoms were excluded.

Tay L B¹⁰⁰et al in their study found that NCS revealed bilateral CTS in 108 (80.6%) and unilateral CTS in 26 (19.4%). There were 35 patients (32.4%) with bilateral CTS being symptomatic unilaterally. Dominant hand involvement was present in 124 patients (92.3%).

In recent publications, by Jablecki CK et al & Padua L et al, the incidence of bilateral symptoms has been reported to be between 60% and 87%,^{104,105}

A. E. Bagatur et al¹⁰⁶ reported the incidence of bilateral symptoms as 59% at presentation and found in their group with unilateral symptoms, more than half had positive electrodiagnostic test results in the asymptomatic, contralateral hand.

Likewise, Corwin and Kasdan¹⁰⁷ using electrodiagnosis, reported that median neuropathy at the wrist may occur in asymptomatic individuals.

Bendler et al¹⁰⁸ also found that 38% of the patients in their series who complained only of unilateral symptoms were shown to have bilateral neurophysiological impairment of the median nerve.

Padua et al¹⁰⁵ found similar circumstances in about half of the asymptomatic hands.

Our study is also consistent with the above mentioned findings in various studies that NCS can detect subclinical CTS in asymptomatic hands.

ETIOLOGY

IDIOPATHIC:

Majority of patients (44.8%) had no obvious cause, were considered to be idiopathic. 35.7% of males, and 48.5% of females had no obvious cause. Nearly two thirds (62.7%) of the idiopathic carpal tunnel syndrome patients had mild to moderate CTS. Almost 14% of idiopathic patients had severe to extreme CTS. Most of the idiopathic CTS (82%) patients had severity of CTS more in their dominant hands.

Reinstein L. et al¹¹¹ have stressed that 'idiopathic CTS' is a bilateral disorder and that it occurs slightly more frequently in the dominant hand.

DIABETES:

Diabetes was present in 13.5% of patient in our study. Diabetes was the 2nd common cause in males (28.6%) and present in 7.4% of the females in our study. Of the 13 patient with diabetes, 3 males and 1 female presented with polyneuropathy. Nearly two third (66%) of patients with diabetes had mild to moderate CTS. Nearly one third (31%) of patients with diabetes had severe to extreme CTS. In the 4 patients with

diabetic polyneuropathy, 2 had severe CTS, 1 had moderate CTS and 1 had mild CTS.

In the study by Urkude R, et al¹⁰⁰ diabetes mellitus was present in 13.4% patients with CTS. Diabetes mellitus is a very commonly occurring systemic disease that is also associated with the presence of CTS. Carpal Tunnel Syndrome develops in 6% of all Type 2 Diabetes sufferers. Median nerve compression is only one of its many complications.¹⁰⁹ According to Dellon et al¹¹⁰, in diabetic patients, the median nerve, as in their other peripheral nerves, is already involved by a polyneuropathy, and is more easily subject to compression.

RHEUMATOID ARTHRITIS:

In our study all the 18 rheumatoid arthritis patients were females and all had carpal tunnel syndrome. Rheumatoid arthritis was the 2nd common cause of CTS among females (26.5%). Most of the patients (88.8%) with rheumatoid arthritis had minimal to moderate CTS, while a few (11.2%) had severe to extreme CTS.

Rheumatoid arthritis is one of the causes that constantly produce a tenovaginitis and consequently CTS.

Polley and Lipscomb (1966) reported that 100 of a series of 1215 with CTS had coexistent rheumatoid arthritis.

Carpal tunnel syndrome (CTS) in patients with RA was investigated by M. Aluclu, et al¹¹² who found that out of 40 adult patients (totally eighty hands) with RA, carpal tunnel syndrome was determined in 20 hands (25%) of the patients with RA. They recommend that an electro physiologic examination should be performed in all

patients with RA as a routine diagnostic procedure.

HYPOTHYROIDISM

In our study, hypothyroidism was seen in 7.3% patients with CTS. Most of them had minimal to moderate CTS.

Thyroid diseases such as hypothyroidism and hyperthyroidism can cause CTS. In hypothyroidism, the swelling of the tissue inside the carpal canal causes the syndrome. CTS can be the first sign of hypothyroidism (Golding DN¹¹³). The substitute medical therapy will improve the clinical symptoms (Fincham RW¹¹⁴). In the study by Stevens JC et al¹⁰² 5.2% of patients with CTS had hypothyroidism at the time of presentation

POLYNEUROPATHY

In our study, apart from 4 patients with diabetic poly neuropathy, 6 patients had polyneuropathy for whom a cause was not established. Polyneuropathy was the third common cause of CTS in males (14.3%) in our study. More than 80% patients with polyneuropathy had moderate to severe CTS.

It is common knowledge that the patients with polyneuropathies are more susceptible to nerve compressions (Dawson DM et al ¹¹⁵). Guillain-Barré also has an association with CTS, but this is not a constant rule (Lambert EH et al¹¹⁶). This also holds true for alcohol-induced neuropathies, where the feet are the extremities that are afflicted first and later on the hands. The above studies show polyneuropathies other than diabetes can present with CTS.

PREGNANCY

Out of 3 pregnant females, 1 was a primigravida with pregnancy induced hypertension (PIH) and other 2 were multigravida without any comorbid illness. All the 3 pregnant patients were in their third trimester. Of 3 pregnant women with CTS 1 had bilateral minimal CTS and 2 had bilateral moderate CTS. All the 3 patients did not come after their deliveries for follow up.

According to the studies done by Bahrami MH et al, Sax TW et al, and Finsen V et al, carpal tunnel Syndrome is common in pregnant women ²⁰⁻²³.

Turgut F et al and Mondelli M et al, in their studies found that, CTS is commonly diagnosed during third trimester of pregnancy and it is often bilateral. In the majority of patients symptoms will resolve either spontaneously or will respond to conservative treatment after delivery. ^{20, 24, 25}.

Mondelli et al¹¹⁷ in their study on CTS in pregnant women found that, CTS in pregnancy is less severe than idiopathic CTS, and has a better spontaneous course.

TRAUMA

In our study, of the 3 patients with trauma, 1 had healed Colles' fracture right wrist and 2 had history of trauma wrists in their symptomatic hands.

CTS can develop as a late onset sequel due to a wrist deformity caused by a carpal bone or radial fracture or other wrist traumas. (Abbott LC) ¹¹⁸. Altimissi et al ¹¹⁹ reported an incidence of CTS in 31% of patients following a Colles fracture.

SENSITIVITY OF VARIOUS ELECTROPHYSIOLOGICAL STUDIES

In our study, of the various electrophysiological tests done, digit 4 sensory median ulnar comparison study was most sensitive (92.1%) followed by F wave comparison studies (87.2%), and the terminal latency index was least sensitive (54.8%). Digit 2 sensory NCS and median motor NCS had a sensitivity of 79.9% and 60.3% respectively. 2nd lumbrical/interossei had a sensitivity of 64.6%.

This is consistent with the sensitivities of Report of the American Association of Electrodiagnostic Medicine, American Academy of Neurology, and the American Academy of Physical Medicine and Rehabilitation (C K Jablecki et al¹²⁰), except that our study showed a greater sensitivity for F wave comparison studies than that of Jablecki et al.

Tests	Present study	C K Jablecki et al
Digit 4 median ulnar	92.1%	85%
Digit 2 sensory	79.9%	65%
Median motor	60.3%	63%
F wave comparison	87.2%	Low sensitivity
Terminal latency index	54.8%	62%
2 nd lumbrical/interossei	64.6%	56%

ULTRASONOGRAM BOTH WRISTS

In our study, ultrasonogram showed evidence of increase in cross sectional area of median nerve in 59 of 93 (63.4%) patients with CTS. In addition to median nerve

thickening, it also showed swelling of tendon sheaths in 12 female patients with rheumatoid arthritis.

Shiu Man Wong, et al¹²¹ studied the usefulness of sonography in a prospective cohort of 133 patients with symptoms suggestive of CTS and found in their study that ultrasonogram had a sensitivity of 83%-94% in detecting CTS.

ElMiedany¹²² compared the results of US with NCS in a group of patients with CTS against a control group and observed a high degree of correlation between the US findings and NCS in diagnosing and in assessing the severity of CTS.

Sharma M et al from Chandigarh, have studied the role of ultrasonography in diagnosis of carpal tunnel syndrome, and found that there was high degree of correlation between ultrasonography and electrophysiological parameters in patients with CTS.

Sonography measures a different parameter (structural pathologic abnormalities of nerve swelling) from that (physiological malfunctions of the median nerve) measured at an electrodiagnostic study. If sonography yields positive results, CTS is confirmed; if the results are negative, one could then refer the patient for an electrodiagnostic study.

Our study had a lower sensitivity when compared to the above studies, which could be due to the interobserver variability and use of poor resolution ultrasound in earlier part of the study which was rectified later by the use of good resolution ultrasound machine and studies were done by a single radiologist.

SUMMARY

SUMMARY

Salient Features in this study were

- ❖ Maximum numbers of patients were in the age group between 40 and 70 years. Females (70.8%) were affected more than males (29.2%). As the duration of symptoms increases there was a progression of severity from mild to extreme CTS.,
- ❖ Paresthesias were seen in more than two third (70.8%) of patients. Tinel's sign was positive in nearly one third of males (32.1%) and females (29.2%). Phalen's test was positive in nearly two third of males (64.3%) and females (64.7%).
- ❖ 19 out of 41 (46.3%) patients with unilateral symptoms showed subclinical CTS in the other hand also after nerve conduction studies.
- ❖ The common causes of carpal tunnel syndrome were idiopathic (44.8%), rheumatoid arthritis (18.8%) diabetes (13.5%) and hypothyroidism (7.3%). Polyneuropathy was the third common cause of CTS in males (14.3%). More than 80% of patients with polyneuropathy had moderate to severe CTS.
- ❖ Of the various electrophysiological tests done, digit 4 sensory median ulnar comparison study was most sensitive (92.1%) followed by F wave comparison studies (87.2%),and the terminal latency index(TLI) was least sensitive.
- ❖ Ultrasonography had a lower sensitivity in detecting carpal tunnel syndrome.

CONCLUSION

CONCLUSION

- ❖ Duration of symptoms with carpal tunnel syndrome correlated very well with the severity in this study.
- ❖ Electrophysiological studies identified subclinical carpal tunnel syndrome in asymptomatic hands in three fourths of patients.
- ❖ Electrophysiological studies helped to assess the severity of carpal tunnel syndrome which could influence the management.

BIBLIOGRAPHY

BIBLIOGRAPHY

1. Eversmann WW Jr. Entrapment and compression neuropathies. In: Green DP, editor. Operative hand surgery vol 2. New York: Churchill Livingstone; 1993. pp 1341 - 85
2. Omer GE, Jr. Median nerve compression at the wrist. *Hand Clin* 1992;**8(2)**:317-24.
3. Blanc PD, Faucett J, Kennedy JJ, Cisternas M, Yelin E. Self-reported carpal tunnel syndrome: predictors of work disability from the National Health Interview Survey Occupational Health Supplement. *Am J Ind Med* 1996;**30(3)**:362-8.
4. Patterson JD, Simmons BP. Outcomes assessment in carpal tunnel syndrome. *Hand Clin* 2002;**18(2)**:359-63, viii.
5. Katz JN, Simmons BP. Clinical practice. Carpal tunnel syndrome. *N Engl J Med* 2002; **346(23)**:1807-12.
6. Tanaka S, Wild DK, Seligman PJ, Behrens V, Cameron L, Putz- Anderson V. The US prevalence of self-reported carpal tunnel syndrome: 1988 National Health Interview Survey data. *Am J Public Health* 1994;**84(11)**:1846-8.
7. Palmer DH, Hanrahan LP. Social and economic costs of carpal tunnel surgery. *Instr Course Lect* 1995;**44**:167-72.

8. Burke FD. Carpal tunnel syndrome: reconciling “demand management” with clinical need. *J Hand Surg [Br]* 2000;**25(2)**:121-7.
9. Atroshi I, Gummesson C, Johnsson R, Ornstein E, Ranstam J, Rosen I. Prevalence of carpal tunnel syndrome in a general population. *JAMA* 1999;**282(2)**:153-8.
10. de Krom MC, Knipschild PG, Kester AD, Thijs CT, Boekkooi PF, Spaans F. Carpal tunnel syndrome: prevalence in the general population. *J Clin Epidemiol* 1992;**45(4)**:373-6.
11. Stevens JC, Sun S, Beard CM, O’Fallon WM, Kurland LT. Carpal tunnel syndrome in Rochester, Minnesota, 1961 to 1980. *Neurology* 1988;**38(1)**:134-8.
12. Einhorn N, Leddy JP. Pitfalls of endoscopic carpal tunnel release. *Orthop Clin North Am* 1996;**27(2)**:373-80.
13. Ferry S, Pritchard T, Keenan J, Croft P, Silman AJ. Estimating the prevalence of delayed median nerve conduction in the general population. *Br J Rheumatol* 1998;**37(6)**:630-5.
14. Prick JJ, Blaauw G, Vredeveld JW, Oosterloo SJ. Results of carpal tunnel release. *Eur J Neurol* 2003;**10(6)**:733-6.
15. Phalen GS. The carpal-tunnel syndrome. Seventeen years’ experience in diagnosis and treatment of six hundred fifty-four hands. *J Bone Joint Surg Am* 1966;**48(2)**:211-28.
16. Mondelli M, Giannini F, Giacchi M. Carpal tunnel syndrome incidence in a general population. *Neurology* 2002;**58(2)**:289-94.

17. Bland JD, Rudolfer SM. Clinical surveillance of carpal tunnel syndrome in two areas of the United Kingdom, 1991-2001. *J Neurol Neurosurg Psychiatry* 2003;**74(12)**:1674-9.
18. Bureau of Labor Statistics. News: lost-work time injuries and illnesses: Characteristics and resulting time away from work. United States: *Bureau of Labour Statistics*; 2001. Available from: <http://www.bls.gov/iif/oshwc/osh/case/osnr0015.pdf>. Last accessed November 2007.
19. Melhorn JM. CTD: carpal tunnel syndrome, the facts and myths. *Kans Med* 1994;**95(9)**:189-92.
20. Stolp-Smith KA, Pascoe MK, Ogburn PL Jr. Carpal tunnel syndrome in pregnancy: frequency, severity, and prognosis. *Arch Phys Med Rehabil* 1998;**79(10)**:1285-7.
21. Bahrami MH, Rayegani SM, Fereidouni M, Baghbani M. Prevalence and severity of carpal tunnel syndrome (CTS) during pregnancy. *Electromyogr Clin Neurophysiol* 2005;**45(2)**:123-5.
22. Sax TW, Rosenbaum RB. Neuromuscular disorders in pregnancy. *Muscle & Nerve* 2006; **34(5)**:559-71.
23. Finsen V, Zeitlmann H. Carpal tunnel syndrome during pregnancy. *Scand J Plast Reconstr Surg Hand Surg* 2006;**40(1)**:41-5.
24. Turgut F, Cetinahin M, Turgut M, Bılkbaşı O. The management of carpal tunnel syndrome in pregnancy. *J Clin Neurosci* 2001;**8(4)**:332-4.

25. Mondelli M, Rossi S, Monti E, Aprile I, Caliandro P, Pazzaglia C *et al.* Long term follow-up of carpal tunnel syndrome during pregnancy: a cohort study and review of the literature. *Electromyogr Clin Neurophysiol* 2007;**47(6)**:259-71.
26. Bernard BP, editor. Musculoskeletal disorders and workplace factors. A critical review of epidemiologic evidence for work-related musculoskeletal disorders of the neck, upper extremity, and low back. NIOSH Publication No 97 - 141. U.S. Department of Health and Human Services, Public Health Service Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health;
 1. 1997. Available from: <http://www.cdc.gov/niosh/docs/97-141/97-141pd.html>.
27. Brian WR, Wright AD. Spontaneous compression of both median nerves in the carpal tunnel. *Lancet* 1947; **1**:277-82.
28. Silverstein BA, Fine LJ, Armstrong TJ. Occupational factors and carpal tunnel syndrome. *Am J Ind Med* 1987;**11(3)**:343-58.
29. Chiang HC, Chen SS, Yu HS, Ko YC. The occurrence of carpal tunnel syndrome in frozen food factory employees. *Gaoxiong Yi Xue Ke Xue Za Zhi* 1990;**6(2)**:73-80.
30. Viikari-Juntura E, Silverstein B. Role of physical load factors in carpal tunnel syndrome. *Scand J Work Environ Health* 1999;**25(3)**:163-85.
31. Stetson DS, Silverstein BA, Keyserling WM, Wolfe WA, Albers JW. Hypothesis relating cumulative trauma to the median nerve with subclinical nerve conduction

- deficits. *Am J Ind Med* 1995;**27(2)**:309-10.
32. Armstrong TJ, Chaffin DB. Carpal tunnel syndrome and selected personal attributes. *J Occup Med* 1979;**21(7)**:481-6.
33. Latko WA, Armstrong TJ, Franzblau A, Ulin SS, Werner RA, Albers JW. Cross-sectional study of the relationship between repetitive work and the prevalence of upper limb musculoskeletal disorders. *Am J Ind Med* 1999;**36(2)**:248-59.
34. Werner R, Armstrong TJ, Bir C, Aylard MK. Intracarpal canal pressures: the role of finger, hand, wrist and forearm position. *Clin Biomech (Bristol, Avon)* 1997;**12(1)**:44-51.
35. Ham SJ, Kolkman WF, Heeres J, den Boer JA. Changes in the carpal tunnel due to action of the flexor tendons: visualization with magnetic resonance imaging. *J Hand Surg [Am]* 1996; **21(6)**:997-1003.
36. Rempel D, Bach JM, Gordon L, So Y. Effects of forearm pronation/ supination on carpal tunnel pressure. *J Hand Surg [Am]* 1998;**23(1)**:38- 42.
37. Hagberg M, Morgenstern H, Kelsh M. Impact of occupations and job tasks on the prevalence of carpal tunnel syndrome. *Scand J Work Environ Health* 1992;**18(6)** :337-45.
38. Vern Putz-Anderson, Bruce P. Bernard, Susan E. Burt, Libby L. Cole, Cheryl Fairfield-Estill, Lawrence J. Fine, *et al.* Musculoskeletal Disorders and Workplace Factors: A Critical Review of Epidemiologic Evidence for Work-Related Musculoskeletal Disorders of the Neck, Upper Extremity, and Low Back: U.S.

Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health; 1997.

39. Lam N, Thurston A. Association of obesity, gender, age and occupation with carpal tunnel syndrome. *Aust N Z J Surg* 1998;**68(3)**:190-3.
40. Becker J, Nora DB, Gomes I, Stringari FF, Seitensus R, Panosso JS, *et al.* An evaluation of gender, obesity, age and diabetes mellitus as risk factors for carpal tunnel syndrome. *Clin Neurophysiol* 2002;**113(9)**:1429-34.
41. Gerr F, Letz R. Risk factors for carpal tunnel syndrome in industry: blaming the victim? *J Occup Med* 1992;**34(11)**:1117-9
42. Nathan PA, Meadows KD, Doyle LS. Occupation as a risk factor for impaired sensory conduction of the median nerve at the carpal tunnel. *J Hand Surg [Br]* 1988;**13(2)**:167-70.
43. Bland JD. The relationship of obesity, age, and carpal tunnel syndrome: more complex than was thought? *Muscle Nerve* 2005;**32(4)**:527-32.
44. Moghtaderi A, Izadi S, Sharafadinzadeh N. An evaluation of gender, body mass index, wrist circumference and wrist ratio as independent risk factors for carpal tunnel syndrome. *Acta Neurol Scand* 2005;**112(6)**:375-9.
45. Boz C, Ozmenoglu M, Altunayoglu V, Velioglu S, Alioglu Z. Individual risk factors for carpal tunnel syndrome: an evaluation of body mass index, wrist index and hand anthropometric measurements. *Clin Neurol Neurosurg*

2004;**106(4)**:294-9.

46. Garland FC, Garland CF, Doyle EJ, Jr., Balazs LL, Levine R, Pugh WM, *et al.* Carpal tunnel syndrome and occupation in U.S. Navy enlisted personnel. *Arch Environ Health* 1996; **51(5)**:395-407.
47. Nathan PA, Istvan JA, Meadows KD. A longitudinal study of predictors of research-defined carpal tunnel syndrome in industrial workers: findings at 17 years. *J Hand Surg [Br]* 2005; **30(6)**:593-8.
48. Spielholz P, Silverstein B, Morgan M, Checkoway H, Kaufman J. Comparison of self-report, video observation and direct measurement methods for upper extremity musculoskeletal disorder physical risk factors. *Ergonomics* 2001;**44(6)**:588-613.
49. Stetson DS, Silverstein BA, Keyserling WM, Wolfe RA, Albers JW. Median sensory distal amplitude and latency: comparisons between non-exposed managerial/professional employees and industrial workers. *Am J Ind Med* 1993;**24(2)**:175-89.
50. Osorio AM, Ames RG, Jones J, Castorina J, Rempel D, Estrin W, *et al.* Carpal tunnel syndrome among grocery store workers. *Am J Ind Med* 1994;**25(2)**:229-45.
51. Schottland JR, Kirschberg GJ, Fillingim R, Davis VP, Hogg F. Median nerve latencies in poultry processing workers: an approach to resolving the role of industrial “cumulative trauma” in the development of carpal tunnel syndrome. *J Occup Med* 1991;**33(5)**:627-31.

52. Moore JS, Garg A. Upper extremity disorders in a pork processing plant: relationships between job risk factors and morbidity. *Am Ind Hyg Assoc J* 1994;**55(8)**:703-15.
53. Palmer KT, Harris EC, Coggon D. Carpal tunnel syndrome and its relation to occupation: a systematic literature review. *Occup Med (Lond)* 2007;**57(1)**:57-66.
54. Seiler JG, 3rd, Milek MA, Carpenter GK, Swiontkowski MF. Intraoperative assessment of median nerve blood flow during carpal tunnel release with laser Doppler flowmetry. *J Hand Surg [Am]* 1989;**14(6)**:986-91.
55. Kiernan MC, Mogyoros I, Burke D. Conduction block in carpal tunnel syndrome. *Brain* 1999;**122(5)**:933-41.
56. Tucci MA, Barbieri RA, Freeland AE. Biochemical and histological analysis of the flexor tenosynovium in patients with carpal tunnel syndrome. *Biomed Sci Instrum* 1997;**33**:246-51.
57. Lundborg G, Dahlin LB, Danielsen N, Hansson HA, Necking LE, Pyykko I. Intra-neural edema following exposure to vibration. *Scand J Work Environ Health* 1987;**13(4)**:326-9.
58. Szabo RM, Slater RR, Jr., Farver TB, Stanton DB, Sharman WK. The value of diagnostic testing in carpal tunnel syndrome. *J Hand Surg [Am]* 1999;**24(4)**:704-14.
59. Buch-Jaeger N, Foucher G. Correlation of clinical signs with nerve conduction tests in the diagnosis of carpal tunnel syndrome. *J Hand Surg [Br]*

1994;**19(6)**:720-4.

60. Katz JN, Larson MG, Sabra A, Krarup C, Stirrat CR, Sethi R, *et al.* The carpal tunnel syndrome: diagnostic utility of the history and physical examination findings. *Ann Intern Med* 1990;**112(5)**:321-7.
61. Gupta SK, Benstead TJ. Symptoms experienced by patients with carpal tunnel syndrome. *Can J Neurol Sci* 1997;**24(4)**:338-42.
62. Kendall WW. Results of treatment of severe carpal tunnel syndrome without internal neurolysis of the median nerve. *J Bone Joint Surg Am* 1988;**70(1)**:151.
63. Yamaguchi DM, Lipscomb PR, Soule EH. Carpal Tunnel Syndrome. *Minn Med* 1965; **48**:22-33.
64. Hoffmann P, Buck-Gramcko D, Lubahn JD. The Hoffmann-Tinel sign. 1915. *J Hand Surg [Br]* 1993;**18(6)**:800-5.
65. Kuhlman KA, Hennessey WJ. Sensitivity and specificity of carpal tunnel syndrome signs. *Am J Phys Med Rehabil* 1997;**76(6)**:451-7.
66. Gelmers HJ. The significance of Tinel's sign in the diagnosis of carpal tunnel syndrome. *Acta Neurochir (Wien)* 1979; **49(3-4)**:255-8.
67. Kushner SH, Ebramzadeh E, Johnson D, Brien WW, Sherman R. Tinel's sign and Phalen's test in carpal tunnel syndrome. *Orthopedics* 1992;**15(11)**:1297-302.
68. Gerr F, Letz R, Harris-Abbott D, Hopkins LC. Sensitivity and specificity of vibrometry for detection of carpal tunnel syndrome. *J Occup Environ Med* 1995;

37(9):1108-15.

69. Golding DN, Rose DM, Selvarajah K. Clinical tests for carpal tunnel syndrome: an evaluation. *Br J Rheumatol* 1986;**25(4)**:388-90.
70. Heller L, Ring H, Costeff H, Solzi P. Evaluation of Tinel's and Phalen's signs in diagnosis of the carpal tunnel syndrome. *Eur Neurol* 1986;**25(1)**:40-2.
71. Mondelli M, Passero S, Giannini F. Provocative tests in different stages of carpal tunnel syndrome. *Clin Neurol Neurosurg* 2001;**103(3)**:178- 83.
72. Burke DT, Burke MA, Bell R, Stewart GW, Mehdi RS, Kim HJ. Subjective swelling: a new sign for carpal tunnel syndrome. *Am J Phys Med Rehabil* 1999;**78(6)**:504-8.
73. De Smet L, Steenwerckx A, Van den Bogaert G, Cnudde P, Fabry G. Value of clinical provocative tests in carpal tunnel syndrome. *Acta Orthop Belg* 1995;**61(3)**:177-82.
74. Kaufman MA. Differential diagnosis and pitfalls in electrodiagnostic studies and special tests for diagnosing compressive neuropathies. *Orthop Clin North Am* 1996;**27(2)**:245-52.
75. Bruske J, Bednarski M, Grzelec H, Zyluk A. The usefulness of the Phalen test and the Hoffmann-Tinel sign in the diagnosis of carpal tunnel syndrome. *Acta Orthop Belg* 2002; **68(2)**:141-5.
76. Katz JN, Stirrat CR, Larson MG, Fossel AH, Eaton HM, Liang MH. A selfadministered hand symptom diagram for the diagnosis and epidemiologic

- study of carpal tunnel syndrome. *J Rheumatol* 1990;**17(11)**:1495-8.
77. Katz JN, Stirrat CR. A self-administered hand diagram for the diagnosis of carpal tunnel syndrome. *J Hand Surg [Am]* 1990;**15(2)**:360-3.
78. Radecki P. A gender specific wrist ratio and the likelihood of a median nerve abnormality at the carpal tunnel. *Am J Phys Med Rehabil* 1994;**73(3)**:157-62.
79. LaBan MM, Friedman NA, Zemenick GA. "Tethered" median nerve stress test in chronic carpal tunnel syndrome. *Arch Phys Med Rehabil* 1986;**67(11)**:803-4.
80. LaBan MM, MacKenzie JR, Zemenick GA. Anatomic observations in carpal tunnel syndrome as they relate to the tethered median nerve stress test. *Arch Phys Med Rehabil* 1989;**70(1)**:44-6.
81. Raudino F. Tethered median nerve stress test in the diagnosis of carpal tunnel syndrome. *Electromyogr Clin Neurophysiol* 2000;**40(1)**:57-60.
82. Stevens JC. AAEE minimonograph #26: The electrodiagnosis of carpal tunnel syndrome. *Muscle Nerve* 1987;**10(2)**:99-113.
83. Rempel D, Evanoff B, Amadio PC, de Krom M, Franklin G, Franzblau A, *et al.* Consensus criteria for the classification of carpal tunnel syndrome in epidemiologic studies. *Am J Public Health* 1998;**88(10)**:1447-51.
84. Practice parameter for electro diagnostic studies in carpal tunnel syndrome (summary statement). American Academy of Neurology, American Association of Electro diagnostic Medicine, and American Academy of Physical Medicine and Rehabilitation. *Neurology* 1993;**43(11)**:2404-5.

85. Practice parameter for carpal tunnel syndrome (summary statement). Report of the Quality Standards Subcommittee of the American Academy of Neurology. *Neurology* 1993;**43**(11):2406-9.

86. Jablecki CK, Andary MT, So YT, Wilkins DE, Williams FH. Literature review of the usefulness of nerve conduction studies and electromyography for the evaluation of patients with carpal tunnel syndrome. AAEM Quality Assurance Committee. *Muscle Nerve* 1993;**16(12)**:1392-414.
87. Chang MH, Liu LH, Lee YC, Wei SJ, Chiang HL, Hsieh PF. Comparison of sensitivity of transcarpal median motor conduction velocity and conventional conduction techniques in electro diagnosis of carpal tunnel syndrome. *Clin Neurophysiol* 2006;**117(5)**:984-91.
88. Dawson DM. Entrapment neuropathies: clinical overview. *Hosp Pract (Minneap)* 1995;**30(8)**:37-40, 43-4.
89. Spinner RJ, Bachman JW, Amadio PC. The many faces of carpal tunnel syndrome. *Mayo Clin Proc* 1989;**64(7)**:829-36.
90. Haig AJ, Tzeng HM, LeBreck DB. The value of electro diagnostic consultation for patients with upper extremity nerve complaints: a prospective comparison with the history and physical examination. *Arch Phys Med Rehabil* 1999;**80(10)**:1273-81.
91. Bingham RC, Rosecrance JC, Cook TM. Prevalence of abnormal median nerve conduction in applicants for industrial jobs. *Am J Ind Med* 1996;**30(3)**:355-61.
92. Bodofsky EB, Wu KD, Campellone JV, Greenberg WM, Tomaio AC. A sensitive new median-ulnar technique for diagnosing mild Carpal Tunnel Syndrome. *Electromyogr Clin Neurophysiol* 2005;**45(3)**:139-44.

93. Keles I, Karagulle Kendi AT, Aydin G, Zog SG, Orkun S. Diagnostic precision of ultrasonography in patients with carpal tunnel syndrome. *Am J Phys Med Rehabil* 2005;**84(6)**:443-50.
94. Koyuncuoglu HR, Kutluhan S, Yesildag A, Oyar O, Guler K, Ozden A. The value of ultrasonographic measurement in carpal tunnel syndrome in patients with negative electro diagnostic tests. *Eur J Radiol* 2005;**56(3)**:365-9.
95. El Miedany YM, Aty SA, Ashour S. Ultrasonography versus nerve conduction study in patients with carpal tunnel syndrome: substantive or complementary tests? *Rheumatology (Oxford)* 2004;**43(7)**:887-95.
96. Mondelli M, Giannini F, Giacchi M. Carpal tunnel syndrome incidence in general population. *Neurology* 2002; 58:289-94.
97. Phalen GS: Reflections of 21 years experience with the carpal tunnel syndrome. *JAMA* 1970; 212:1365-1367
98. Radecki P :The familial occurrence of carpal tunnel syndrome. *Muscle Nerve* 1994;17:325-330.
99. Bland JD, Rudolfer SM. Clinical surveillance of carpal tunnel syndrome in two areas of the United Kingdom, 1991-2001. *J Neurol Neurosurg Psychiatry* 2003;74(12):1674-9.
100. Tay L B, Urkude R, Verma K K. Clinical profile, electrodiagnosis and outcome in patients with carpal tunnel syndrome: a Singapore perspective. *Singapore Med J* 2006; 47(12) : 1049

101. Tanaka S, Wild DK, Seligman PJ, Behrens V, Cameron L, Putz-Anderson V. The US prevalence of self-reported carpal tunnel syndrome: 1988 National Health Interview Survey data. *Am J Public Health* 1994; 84(11):1846-8.
102. Stevens JC, Sun S, Beard CM, O'Fallon WM, Kurland LT. Carpal tunnel syndrome in Rochester, Minnesota, 1961 to 1980. *Neurology* 1988;38(1):134-8.
103. Kendall WW. Results of treatment of severe carpal tunnel syndrome without internal neurolysis of the median nerve. *J Bone Joint Surg Am* 1988;70(1):15
104. Jablecki CK, Andary MT, So YT, Wilkins DE, Williams FH. Literature review of the usefulness of nerve conduction studies and electromyography for the evaluation of patients with carpal tunnel syndrome. AAEM Quality Assurance Committee. *Muscle Nerve* 1993;16:1392-414.
105. Padua L, Padua R, Nazzaro M, Tonali P. Incidence of bilateral symptoms in carpal tunnel syndrome. *J Hand Surg [Br]* 1998;23:603-6
106. A. E. Bagatur, G. Zorer The carpal tunnel syndrome is a bilateral disorder
THE JOURNAL OF BONE AND JOINT SURGERY VOL. 83-B, NO. 5,
JULY 2001
107. Corwin HM, Kasdan ML. Electrodiagnostic reports of median neuropathy at the wrist. *J Hand Surg [Am]* 1998;23:55-7.
108. Bendler EM, Greenspun B, Yu J, Erdman WJ 3rd. The bilaterality of carpal

- tunnel syndrome. *Arch Phys Med Rehabil* 1977;58:362-4.
109. Brown MJ, Asbury AK (1984) Diabetic neuropathy *Ann Neurol* 15: 2–12
110. Dellon AL (1992) Treatment of symptomatic diabetic neuropathy by surgical decompression of multiple peripheral nerves. *Plast Reconstr Surg* 89: 689–697
111. Reinstein L. Hand dominance in CTS. *Arch Phys Med Rehab* 1981; 62: 202–203.
112. M. Aluclu, A. Turhanoglu & M. Aluclu : The Frequency Of Carpal Tunnel Syndrome In Patients With Rheumatoid Arthritis . *The Internet Journal of Neurology*. 2006 Volume 5 Number 2
113. Golding DN (1970) Hypothyroidism presenting with musculoskeletal symptoms. *Ann Rheum Dis* 29: 10–14
114. Fincham RW, Cape CA (1968) Neuropathy in myxedema. A study of sensory nerve conduction in the upper extremities. *Arch Neurol* 19: 464–466
115. Dawson DM, Hallett M, Millender LH (1983) Entrapment neuropathies. Boston, Little Brown Co.
116. Lambert EH, Mulder DW (1964) Nerve conduction in the Guillain-Barré syndrome. *Electroencephalogr Clin Neurophysiol* 17: 86
117. Mondelli, Stefania Rossi. Prospective study of positive factors for improvement of carpal tunnel syndrome in pregnant women. *Muscle & Nerve* 2007; 36:778-783
118. Abbott LC, Saunders JB deCM(1933) Injuries of the median nerve in fractures of the lower end of the radius. *Surg Gynecol Obstet* 57: 507–516

119. Altissimi M, Antenucci R, Fiacca C, Mancini GB (1986) Long-term results of conservative treatment of fractures of the distal radius. *Clin Orthop* 206: 202–210
120. C. K. Jablecki, M. T. Andary, M. K. Floeter, R. G. Miller, C. A. Quartly, M. J. Vennix and J. R. Wilson. Practice parameter: Electrodiagnostic studies in carpal tunnel syndrome: Report of the American Association of Electrodiagnostic Medicine, American Academy of Neurology, and the American Academy of Physical Medicine and Rehabilitation. *Neurology* 2002;58:1589-1592
121. Shiu Man Wong, MD James F. Griffith, MD Andrew C. F. Hui, MD Carpal Tunnel Syndrome: Diagnostic Usefulness of Sonography. *Radiology* 2004; 232:93–99
122. El Miedany YM, Aty SA, Ashour S. Ultrasonography versus nerve conduction study in patients with carpal tunnel syndrome: substantive or complementary tests? *Rheumatology (Oxford)* 2004;43(7):887-95.
123. Sharma M Prabhakar S role of ultrasonography in the diagnosis of carpal tunnel syndrome. *Annals of Indian Academy of Neurology*, volume 10, supplement 2, 2007.
124. Mondelli M, Giannini F, Giacchi M. Carpal tunnel syndrome incidence in general population. *Neurology* 2002; 58:289-94.
125. Bland JD. Do nerve conduction studies predict the outcome of carpal tunnel compression? *Muscle Nerve* 2001; 24:935-40.
126. Mondelli M, Giannini F, Vecchierelli B, et al. Diagnostic pathway in carpal tunnel syndrome. *Riv Neurobiol* 2000; 46:301-5.

127. Padua L, Lo Monaco M, Gregori B, et al. Neurophysiological classification and sensitivity in 500 carpal tunnel syndrome hands. *Acta Neurol Scand* 1997; 96: 211-17.

PROFORMA

PROFORMA

“Evaluation of Clinical, Electrophysiological, and
Radiological Profile of Carpal Tunnel Syndrome”

Institute of Neurology
Madras Medical College, Chennai – 3.

Name:

Age/Sex:

MIN No:

HISTORY:

DURATION:

Paraesthesia - Nocturnal/Diurnal /both
- D1/D2/D3/D4/D5/ Thenar/ Forearm/Arm
- Right/Left

Pain wrists - Right/Left

Sensory loss - D1/D2/D3/D4/D5/ Thenar/ forearm/arm
- Right/ Left

Weakness

Atrophy

Joint swelling

Arthralgia

Pregnancy-Yes /No

Amenorrhoea- MA

Cold intolerance

Hoarseness of voice

Neck swelling

Polyuria/polydipsia/polyphagia

PAST HISTORY:

DM/HT/HYPOTHYROIDISM/RA/TRAUMA/TB/RENAL FAILURE

FAMILY HISTORY:

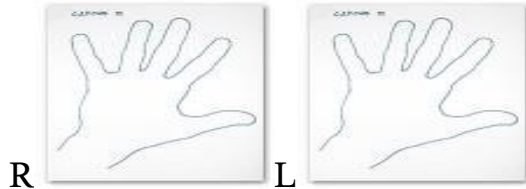
OCCUPATION:

Manual labour/housewife/typist/computer operator/others

PERSONAL HISTORY: smoker/alcoholic/substance abuse

SIGNS:

SENSORY LOSS



WEAKNESS: LUMBRICALS/OPP/APB/FPB/OTHERS

WASTING:

REFLEX LOSS:

TINEL'S SIGN:

PHALEN'S TEST:

GOITRE

DEFORMITY- SWAN NECK/ BUTTON HOLE/ ULNAR DEVIATION

OTHERS:

INVESTIGATIONS:

RA fr	Blood sugar			THYROID(TFT)		
	F	PP	GTT	T3	T4	TSH

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INSTITUTE OF NEUROLOGY
MADRAS MEDICAL COLLEGE, CHENNAI – 3.

*“Evaluation of Clinical, Electrophysiological, and Radiological Profile of
Carpal Tunnel Syndrome”*

NERVE CONDUCTION STUDY

Name:
MIN No:

Age/Sex:
Unit:

MOTOR NERVE CONDUCTION STUDY

NERVE	LATENCY (m sec)		AMPLITUDE (mV)		NCV (m/s)	F wave latency (m sec)
	Distal	Proximal	Distal	Proximal		
MEDIAN R						
ULNAR R						
MEDIAN L						
ULNAR L						
PALM and WRIST STIMULATION COMPARISON MEDIAN R						
PALM and WRIST STIMULATION COMPARISON MEDIAN L						
2ND LUMBRICAL/ INTEROSSEI R	M					
	U					

2ND LUMBRICAL/ INTEROSSEI L	M						
	U						

TLI MEDIAN R=

TLI MEDIAN L=

SENSORY NERVE CONDUCTION STUDY

NERVE		LATENCY (milli sec)	AMPLITUDE (µV)	VELOCITY (m/s)
MEDIAN R (DIGIT 2)				
ULNAR R (DIGIT 5)				
MEDIAN L (DIGIT 2)				
ULNAR R (DIGIT 5)				
DIGIT 4 SENSORY R	M			
	U			
DIGIT 4 SENSORY L	M			
	U			
PALM and WRIST STIMULATION COMPARISON MEDIAN R	M			
	U			
PALM and WRIST STIMULATION COMPARISON MEDIAN L	M			
	U			
	R			
	L			
	R			
	L			

	R			
	L			

EMG STUDIES

MUSCLE	IN	SPON	MUP			RECRUITMENT
			AMP	DUR	PHASES	

ELECTROPHYSIOLOGICAL STAGING

ULTRASONOGRAM BOTH WRISTS

USG wrist	
R	L

CONCLUSION: