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### THE EFFECT OF NEURAL MOBILIZATION WITH CERVICAL TRACTION IN CERVICAL RADICULOPATHY PATIENTS

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#### ABSTRACT

Transcutaneous Electrical Nerve Stimulation (TENS) is the application of electrical stimulation to the skin via surface electrodes to stimulate nerve fibers, primarily used for relief of pain. The gate control theory of pain asserts that non-painful input closes the nerve "gates" to painful input, which prevents pain sensation from traveling to the central nervous system. Neurodynamics or Neural mobilization, refers to the communication between different parts of the nervous system and to the nervous systems relationship to the musculoskeletal system. It has been shown that the nerves move independently from other tissues. The term Neurodynamics first made an appearance in 1989 and has since been further developed over the last 30 years. Neurodynamics is now seen as an important part of injury assessment and treatment. Spondylosis, or spinal arthritis, is a medical term for the normal wear and tear that occurs in the joints and bones of the spine as people get older. Cervical spondylosis refers to these age related changes that occur in the neck (the cervical spine). Cervical radiculopathy (CR) is a symptom complex of neck pain and radiating arm pain due to compression of one or more cervical nerve roots. The impingement is caused by spondylotic narrowing of the intervertebral foramina, by intervertebral disc herniation or by both. Cervical radiculopathy typically manifests as pain radiating from the neck into the distribution of the affected root. The exact location and pattern of pain may vary widely, and a classic dermatomal distribution of pain is not always present. Associated sensory, motor, and reflex disturbances may or may not be present. Because acute cervical radiculopathy generally has a self-limited clinical course, nonsurgical treatment is the appropriate initial approach for most patients. Surgical treatment may be considered when nonsurgical treatment fails and in the patient with a significant neurologic deficit. Methods: Experimental study design, 60 subjects with Unilateral Cervical Radiculopathy and ULTT-1 positive for median nerve bias, randomized 20 subjects each into three groups - Group A, B and C respectively. Group A received both Cervical Traction and Neural Mobilization. The Group B received only Mechanical Cervical Traction. The Group C received only Neural Mobilization. The duration of intervention was given 3 treatment sessions per week for four weeks. Outcome measures such as Numerical Pain Rating Scale, Global Rating of Change Scale, and Neck Disability Index were measured before, at the end of 2nd and 4th week post treatment. Conclusion: The present study concludes that simultaneous application of mechanical cervical traction with neural mobilization is more effective in improving pain, functional disability and severity of radicular symptoms than mechanical cervical traction and neural mobilization alone for subjects with unilateral cervical radiculopathy.

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# **INTRODUCTION**

Cervical radiculopathy (CR) is, a disorder of the cervical nerve root (CNR) and most commonly is caused by a cervical disc herniation, spondylotic spur or cervical osteophyte or other space - occupying lesion, resulting in nerve root inflammation, impingement or both which may lead to chronic pain and disability. The average annual incidence rate of cervical radiculopathy is 83 per 100 000 for the population in its entirety, with an increased prevalence occurring in

the fifth decade of life (203 per 100 000). The location and pattern of symptoms in Cervical radiculopathy vary depending on nerve root level affected and primarily presents with sensory symptoms into the upper limb such as pain, numbness, tingling sensation; motor symptoms like muscle weakness; and reflex hypo activity which often result in significant functional limitations and disability. Treatment techniques such as Cervical Traction and Neural Mobilization both have been individually advocated for treatment of Cervical radiculopathy due to their various effects. Neural tissue mobilization techniques (NMTs) theorize to examine the neural

tension in nerves and mobilize the nerves that exhibit neural tension by passive or active movements by using Tensioning, Sliding and Single Joint Movement techniques and focused on restoring the ability of the nervous system to tolerate the normal compressive, friction, and tensile forces associated with daily and sport activities. With this method, tension was gently applied to the involved nerve root that caused mild pulling but no pain and a low amplitude repetitive movement was introduced in the direction of perceived neural tension. NMTs are widely used to normalize the CNR's structure and function by reducing nerve adherence, facilitating nerve gliding and decreasing neural mechanosensitivity in patients with cervical radiculopathy. Mechanical cervical traction is technique used to decompress the nerve root by separating the cervical segments through long-axis traction. Many studies reveals that Intermittent Cervical traction for cervical radiculopathy found to be effective in reduction of pain and disability; however, no standard parameters have been reported. It is speculated that improvements in symptoms are due to distraction of articular surfaces by traction which unloads the components of the spine by stretching muscles, ligaments and functional units. Cervical traction and neural mobilization techniques (NMTs) have been advocated in the management of CR due to their immediate analgesic effect.12-19. The analgesic effect of these two techniques has been explored and recognized in many RCT studies and in systematic reviews. The present study was to find out whether the simultaneous application of Cervical Traction and Neural Mobilization does have effect on improving neck pain, radicular symptoms and neck disability in subjects with unilateral cervical radiculopathy? Hence, the purpose of study with objective is to find the effect of simultaneous application of cervical traction and neural mobilization on improving neck pain, radicular symptoms and neck disability in subjects with Cervical Radiculopathy. It was hypothesised that Simultaneous application of cervical traction and neural mobilization will have a significant difference on improving neck pain, radicular symptoms and neck disability in subjects with unilateral cervical radiculopathy.

## **MATERIAL & METHODOLOGY**

**Study Approval -** The Maharaj Vinayak Global University (MVGU), Department of Research Committee (DRC) served as the approval board for this study. This study was submitted as a full board review, and the board members approved the study. **Study** A repeated measures experimental study design with three groups. Subjects included were both sex with age group between 45 to 55 years , with history of neck pain associated with radicular pain below the occiput and unilateral side of the neck with a tingling sensation occurring on the lateral aspectof the elbow joint, positive for the test item cluster for provocative Tests which include the Spurling's test, Shoulder Abduction test, Valsalva Maneuver, Neck Distraction test and Upper limb tension test, positive history from the Six historical questions diagnostic for Cervical Radiculopathy which include the following questions:

- 1) Which are your most bothersome symptoms: Pain, Numbness / Tingling, Loss of feeling.
- Where are the symptoms mostbothersome: Neck, Shoulder, Scapula, Arm above elbow, Arm below elbow, Hand, Fingers.
- 3) Symptom behaviour Constant, Intermittent, Variable.
- 4) Entire limb numbness.
- 5) Are the symptoms keeping you away from sleep?
- 6) Does the neck movement improve orworsen the arm pain.
- 7) Complaints of pain radiating to Upper Extremity that was provoked or exacerbated by

Cervical Range of Motion, Paraesthesia in dermatomal pattern (for Median Nerve), mild to moderate severity of radicular symptoms measured by Visual Analogue Scale between 5 to 7 cm, subjects having clear or unequivocal relief of the radicular pain with the manual cervical distraction test, and positive for ULTT1, Median Nerve Bias. Subjects excluded were with sensory loss or motor weakness due to cervical radiculopathy, history of trauma, rheumatoid arthritis, malignancy, cervical Instability, cord myelopathy, vertebrobasilar syndrome, undiagnosed pain, who has undergone cervical surgeries, bilateral radiculopathy, neuropathies etc. Total 60 Subject (n=60) who meet the inclusion criteria were recruited by Simple random sampling method using closed envelops, randomly allocated 20 subjects into each Group A, B, and C. Subjects who meet inclusion criteria were informed about the study and a written informed consent was taken. Total four weeks of intervention with 12 treatment sessions which includes 3 treatment sessions in a week was given.

Procedure: Group A: In this group subjects were treated simultaneously with mechanical cervical traction and neural mobilization. Calibrated Biomed Traction unit was used in the study. For Intermittent Cervical Traction subjects were instructed about the procedure and were taken in supine on the treatment couch with the body in neutral position. The cervical spine was placed at an angle of approximately 15 degrees of flexion with traction force to be 10% of subject's body weight and was increased approximately 1 kg every visit, depending on centralization or reduction of symptoms and on/off cycle set at 60/30 seconds) and treatment was given for 9 minutes persession. A buzar switch in the unaffected arm was given so that subject can alert the therapist in case of any discomfort during traction and neural mobilization. After starting the traction machine, neural mobilization procedure was performed. The therapist stood on the affected side besides the subject and depressed the subject's shoulder with one hand while the elbow would be kept in 90 degrees of flexion and forearm in supination and wrist and fingers in extended position using other hand. Thensubject's arm was passively taken into 90 to 100 degrees of abduction (as tolerable by the subject). This was followed by either the sliding or the gliding procedure. If sliding of the nerve was to bedone then alteration of elbow extension (loading median nerve) with wrist flexion (unloading median nerve) and elbow flexion (unloading median nerve) with wrist extension (loading median nerve) was performed for 6 sets of repetitions. Each set was performed in a slow oscillatory manner during the pull time of the traction (60 seconds) followed by 30 seconds rest.23 In the 60 seconds each time first 3 sets of sliding followed by 3 sets of gliding were given, followed by 30 seconds rest. As the symptoms improved the patient was progressed to gliding/ tensioning technique.

**Group B:** In this group subjects were treated with only mechanical cervical traction. The Procedure forgiving traction was followed same as given inGroup A, but no neural mobilization was given along with it.

**Group C:** In this group subjects were treated with only neural mobilization (median nerve). The procedure of applying neural mobilization was the same as given in Group A, but no traction was applied.

Outcome Measures: Outcome measures were measured before at theend of 2<sup>nd</sup> and 4<sup>th</sup> week of treatment. Pain as recorded by NPRS (Numerical Pain Rating Scale): Subjects was asked to mark anywhere on the scale from 0 to 10 which best describes his/ her pain intensity. The mark was recorded using a measuring scale (in cm) to measure the marked point and the obtained NPRS Score was recorded and used for analysis. The Severity of Radicular Symptoms was recorded by the GROC (Global Rating of Change Scale) which is a self administered scale, being filled in by participants themselves. The patient has to rate his/her overall radiculopathy condition from the time that the treatment began until now by marking on any of the numbers from -7 to 7, where it starts from worse i.e. negative to better. The Functional Disability was recorded by NDI (Neck Disability Index) which consists of 10 sections which have Questions that give information as to how the neck pain has affected the ability to manage in everyday life. Each section was scored on a 0 to 5 scale with first statement being 0 (i.e. No pain) and last statement being 5 (i.e. worst imaginable pain). Higher score indicates participant has more disability. The Neck Disability Index and Numeric Pain Rating Scale exhibited fair test-retest reliability, with adequate responsiveness in patient with cervical radiculopathy. The NDI is the most widely used and most strongly validated instrument for assessing self-rated disability in patients with neck pain. Global Rating of Change Scale offers aflexible, quick and simple method of charting self assessed clinical progress in research and clinical settings and has clinical relevance, adequate reproducibility and sensitivity to changes. Test retest reliability was found to be high (ICC 0.9) and face validity (0.90).

**Data Analysis:** Descriptive statistical analysis carried out in the present study. Out Come measurements assessed at 5% level of significance with p-value was set at 0.05 less than this is considered as statisticallysignificant difference. Repeated Measures Analysis of Variance was used for within the group analysis and Bonferroni's as post-hoc test was used to find the significance in pair-wise comparison. One-way ANOVA and Tukey HD post-hoc test and Kruskal-Wallis Test have been used to compare the means of variables between the three groups and multiple pair-wise comparisons with calculation of percentage of difference between the means.

## RESULTS

Total 60 subjects were studied, in Group A there were 20 subjects with mean age 50.55 years, in Group B there were 20 subjects with mean age 49.71 years and in Group C there were 20 subjects with mean age 49.89 years. Within the group analysis of outcome measures in Group A, GroupB, and Group C shows that there is a statistically significant change in means of NPRS and NDI when means were analyzed from pre intervention to  $2^{nd}$  week intervention and to 4 weeks postintervention measurements within the groups.

In Group A the severity of radicular Symptoms was recorded by the GROC scale before intervention there were 35% of subjects had a condition of 'Quite a bit worse', 30% of subjects were 'Moderately worse', 20% of subjects were with 'Somewhat worse' and15% subjects with 'A little bit worse'. In Group A after 2 weeks of intervention 10% of subjects were with condition 'Somewhat worse', 25% of subjects with a 'Alittle bit worse', 15% of subjects with 'A tiny bit worse', 25% of subjects with a 'A tiny bit better', 25% of subjects with 'A little bit better'. After 4 weeks of intervention 15% of subjects with 'A tiny bit worse', 20% of subjects with 'A tiny bit better', 5% of subjects with 'A little bit better' 15% of subjects with 'somewhat better', 20% of subjects with 'Moderately better', 20% of subjects with Quite a bit better, 5% of subjects with 'A great deal better'. In Group B the severity of radicular Symptoms was recorded by the GROC scale before intervention there before intervention there were 40% of subjects had a condition of 'moderate worse', 25% of subjects were 'Somewhat worse', 10% of subjects were 'Alittle bit worse'. In Group B after 2 weeks of intervention 10% of subjects were with condition 'Somewhat worse', 25% of subjects with a 'A little bit worse', 40% of subjects with 'A tiny bit worse', 25% of subjects with a 'A tiny bit better'. After 4 weeks of intervention 10% of subjects with 'A tiny bit worse', 25% of subjects with 'A tiny bit better', 30% of subjects with 'A little bit better', 25% of subjects with 'somewhat better', 10% of subjects with 'Moderately better'. In Group C the severity of radicular Symptoms before intervention there were 25% of subjectshad a condition of 'Quite a bit worse', 30% of subjects were 'Moderately worse', 25% of subjects were with 'Somewhat worse' and 20% subjects with 'A little bit worse'. In Group C after 2 weeks of intervention 20% of subjects were with condition 'Somewhat worse', 35% of subjects with a 'A little bit

Table 1. Analysis of NPRS and NDI within the Group A (Repeated measures analysis)

|                 | Pre intervention  | 2 <sup>nd</sup> week | 4 <sup>th</sup> week |
|-----------------|-------------------|----------------------|----------------------|
| Study Group     | (Mean±SD) min-max | (Mean±S) min-max     | (Mean±S) min-max     |
| NPRS            | $6.15 \pm 0.62$   | 4.26±0.90            | $1.77 \pm 0.65$      |
|                 | (5.2-6.9)         | (2.5-5.8)            | (1.2-3.3)            |
| Neck Disability | 49.41±11.70       | 34.21±7.56           | $19.91 \pm 3.91$     |
| Index in %      | (28-70)           | (18-46)              | (28-70)              |

|                            |   | Percentageof<br>change F-Value | E Valua | Significance  | Effort size r | 95% Confidence Interval for Difference |       |
|----------------------------|---|--------------------------------|---------|---------------|---------------|--|-------|
|                            |   |                                | P-value | Effect size i | Lower Bound   | Upper Bound                            |       |
|                            | Pre to 2 <sup>nd</sup> week             | -31.78%                        | 183.31  | P=0.000       | +0.78 (Large) | 5.84                                   | 6.44  |
| NPRS                       | 2 <sup>nd</sup> to 4 <sup>th</sup> week | -58.58%                        | 234.71  | P=0.000       | +0.85 (Large) | 3.83                                   | 4.68  |
|                            | Pre to 4 <sup>th</sup> week             | -72.33%                        | 972.71  | P=0.000       | +0.97 (Large) | 1.45                                   | 2.08  |
| Neck Disability Index in % | Pre to 2 <sup>nd</sup> week             | -31.76%                        | 104.12  | P=0.000       | +0.62 (Large) | 43.92                                  | 54.86 |
|                            | 2 <sup>nd</sup> to 4 <sup>th</sup> week | -42.81%                        | 104.13  | P=0.000       | +0.77 (Large) | 30.66                                  | 37.75 |
|                            | Pre to 4 <sup>th</sup> week             | -58.71%                        | 154.89  | P = 0.000     | +0.87 (Large) | 18.06                                  | 21.74 |

Table 2. Analysis of NPRS and NDI within the Group B (Repeated measures analysis)

| Study Group     | Pre intervention  | 2 <sup>nd</sup> week | 4 <sup>th</sup> week |  |
|-----------------|-------------------|----------------------|----------------------|--|
|                 | (Mean±SD) min-max | (Mean±SD) min-max    | (Mean±SD) min-max    |  |
| NPRS            | 6.28±0.52         | 4.79±0.94            | 2.83±0.88            |  |
|                 | (5.2-6.8)         | (2.6-6.3)            | (1.1-4.3)            |  |
| Neck Disability | 53.20±12.37       | 36.90±9.30           | 25.00±6.60           |  |
| Index in %      | (30-76)           | (20-52)              | (14-38)              |  |

|            |   | Percentageof E-value |                | Significance | Effort size r | 95% ConfidenceInterval for Difference |             |
|------------|---|----------------------|----------------|--------------|---------------|---------------------------------------|-------------|
|            |   | change               | change F-value |              | Effect size i | Lower Bound                           | Upper Bound |
|            | Pre to 2 <sup>nd</sup> week             | -24.72%              | 109.02         | P = 0.000    | +0.71 (Large) | 1.07                                  | 1.92        |
| NPRS       | 2 <sup>nd</sup> to 4 <sup>th</sup> week | -41.91%              | 152.89         | P = 0.000    | +0.74 (Large) | 1.49                                  | 2.42        |
|            | Pre to 4 <sup>th</sup> week             | -55.93%              | 536.53         | P = 0.000    | +0.93 (Large) | 3.01                                  | 3.87        |
| Neck       | Pre to 2 <sup>nd</sup> week             | -31.63%              | 182.17         | P = 0.000    | +0.58 (Large) | 12.74                                 | 19.85       |
| Disability | 2 <sup>nd</sup> to4 <sup>th</sup> week  | -33.24%              | 158.36         | P = 0.000    | +0.58 (Large) | 9.11                                  | 14.68       |
| Index in % | Pre to 4 <sup>th</sup> week             | -54.00%              | 239.22         | P = 0.000    | +0.82 (Large) | 22.84                                 | 33.57       |

There is a clinical significance effect with large effect size. Analysis of severity of radicular Symptoms between the groups found that GroupA shown greater percentage of reduction inseverity of radicular Symptoms that Group B and C. worse', 25% of subjects with 'A tiny bit worse', 20% of subjects with a 'A tiny bit better'. After 4 weeks of intervention 15% of subjects with 'A tiny bit worse', 20% of subjects with 'A tiny bit better', 20% of subjects with 'A little bit better', 35% of subjects with 'somewhat better', 10% of subjects with 'Moderately better'.

| Table 3. Analysis of NPRS and NDI within the Group C (Repeated measures analysis | s) |
|--|----|
|--|----|

| Study Group     | Pre intervention (Mean±SD) | 2 <sup>nd</sup> week (Mean±SD) | 4 <sup>th</sup> week (Mean±SD) |  |
|-----------------|----------------------------|--------------------------------|--------------------------------|--|
|                 | min-max                    | min-max                        | min-max                        |  |
| NPRS            | 6.15±0.62                  | 4.70±0.89                      | 2.45±1.03                      |  |
|                 | (5.1-6.8)                  | (3.1-5.9)                      | (1.1-4.3)                      |  |
| Neck Disability | 50.80±11.27                | 37.60±8.77                     | 23.80±6.67                     |  |
| Index in %      | (28-70)                    | (20-54)                        | (14-38)                        |  |

|  |   | Percentage of | F-value | Significance | Effect size r | 95% Confidence Interval for<br>Difference |             |
|--|---|---------------|---------|--------------|---------------|---|-------------|
|  |   | change        |         | P-value      |               | Lower Bound                               | Upper Bound |
| NPRS Pre to $2^{nd}$<br>Pre to $4^{th}$<br>Pre to $4^{th}$ | Pre to 2 <sup>nd</sup> week             | -23.58%       | 191.22  | P=0.000      | +0.69 (Large) | 5.87                                      | 6.44        |
|  | 2nd to 4 <sup>th</sup> week             | -47.88%       | 305.83  | P=0.000      | +0.76 (Large) | 4.28                                      | 5.12        |
|  | Pre to 4 <sup>th</sup> week             | -60.17%       | 480.84  | P=0.000      | +0.91 (Large) | 1.97                                      | 2.94        |
| Neck Disability  | Pre to 2 <sup>nd</sup> week             | -25.97%       | 233.78  | P=0.000      | +0.54 (Large) | 45.52                                     | 56.09       |
|  | 2 <sup>nd</sup> to 4 <sup>th</sup> week | -32.25%       | 158.98  | P=0.000      | +0.66 (Large) | 33.48                                     | 41.71       |
|  | Pre to 4 <sup>th</sup> week             | -53.15%       | 255.54  | P=0.000      | +0.82(Large)  | 20.1                                      | 26.88       |

#### Table 4: Comparison of NPRS score between the three groups- Group A, B and C

|  | Percentageof<br>difference | Effect size r | Between two Groups<br>Significance | 95%Confidence interval<br>of the difference |        | Between<br>Three Groups |
|--|----------------------------|---------------|------------------------------------|---|--------|-------------------------|
|  |                            |               | P-value                            | Lower                                       | Upper  |                         |
| Pre intervention NPRS Score                  |                            |               |                                    |   |        |                         |
| Group A & B                                  | 2.26%                      | 0.12 Small    | P=0.733 (NS)                       | -0.587                                      | 0.308  | 0.396                   |
| Group B & C                                  | -2.08%                     | 0.12 Small    | P=0.763 (NS)                       | -0.316                                      | 0.578  | p=0.821 (NS)            |
| Group A & C                                  | 0.16%                      | +0.00 Small   | P=0.997 (NS)                       | -0.457                                      | 0.438  |                         |
| 2 <sup>nd</sup> week of intervention         |                            |               |                                    |   |        |                         |
| Group A & B                                  | -11.95%                    | +0.27 Small   | P=0.1577 (NS)                      | -0.468                                      | 0.156  | 4.002                   |
| Group B & C                                  | -1.88%                     | +0.06 Small   | P=0.954 (NS)                       | -0.447                                      | 0.776  | p=0.1356(NS)            |
| Group A & C                                  | 10.06%                     | +0.024 Small  | P=0.266 (NS)                       | -1.141                                      | 0.241  |                         |
| 4 <sup>th</sup> Post Intervention NPRS Score |                            |               |                                    |   |        |                         |
| Group A & B                                  | 46.63%                     | +0.56 (Large) | P=0.001                            | -1.718                                      | 412    | 12.936                  |
| Group B & C                                  | -14.38%                    | +0.18 (Small) | P=0.357 (NS)                       | -0.278                                      | 1.028  | p=0.002                 |
| Group A & C                                  | 32.86%                     | +0.38(Medium) | P=0.037                            | -1.345                                      | -0.037 |                         |

| Table 5. Comparison of NDI between the Groups A, B and | С |
|--|---|
|--|---|

|                                      | Percentageof<br>difference | Effect size r  | Between two<br>Groups<br>Significance | 95%Confidence interval<br>of the difference |       | Between<br>Three<br>Groups |
|--------------------------------------|----------------------------|----------------|---------------------------------------|---|-------|----------------------------|
|                                      |                            |                | P-value                               | Lower                                       | Upper | 1                          |
| Pre intervention NDI                 |                            |                |                                       |   |       |                            |
| Group A & B                          | 7.41%                      | 0.16 Small     | P=0.567 (NS)                          | -12.78                                      | 5.16  | F=0.532                    |
| Group B & C                          | -62.31%                    | 0.11 Small     | P=0.798 (NS)                          | -6.58                                       | 11.36 | p=0.592                    |
| Group A & C                          | 2.78%                      | +0.07 Small    | P=0.924 (NS)                          | -10.38                                      | 7.56  | (NS)                       |
| 2 <sup>nd</sup> week of intervention |                            |                |                                       |   |       |                            |
| Group A & B                          | 7.58%                      | +0.16 Small    | P=0.584 (NS)                          | -9.24                                       | 3.84  | F=0.877                    |
| Group B & C                          | 1.88%                      | +0.04 Small    | P=0.965 (NS)                          | -7.24                                       | 5.84  | p=0.423                    |
| Group A & C                          | 9.48%                      | +0.21 Small    | P=0.426 (NS)                          | -9.94                                       | 3.14  | (NS)                       |
| 4 <sup>th</sup> Post Intervention    |                            |                |                                       |   |       |                            |
| Group A & B                          | 22.72%                     | +0.43 (Medium) | P=0.022                               | -9.56                                       | 66    | E-4 152                    |
| Group B & C                          | -4.92%                     | +0.07 (Small)  | P=0.794(N)                            | -3.24                                       | 5.66  | r = 4.155<br>r = 0.022     |
| Group A & C                          | 17.85%                     | +0.34 (Medium) | P=0.098                               | -8.36                                       | 0.56  | p=0.022                    |

The NPRS and NDI means were compared between the three groups. Before intervention shows that there was no statistically significant difference between the groups at pre-intervention and  $2^{nd}$  week post intervention. When the means were compared at  $4^{th}$  week post-intervention shown that there was a statistically significant difference between the groups A & B, and Group A & C and there is no statistically significant difference between the Group B & C.

## DISCUSSION

It is found that 4 weeks of interventions consisting simultaneous application of Mechanical Cervical Traction along with Neural Mobilization for the subjects in Group A shown statistically significant greater percentage of improvement in pain, functional disability and severity of the radicular symptoms than Group B who received only Mechanical Cervical Traction and Group C who received only NeuralMobilization.

In Group A subjects the improvements could be because of simultaneous application of Mechanical Cervical Traction along with Neural Mobilization. In this present study maintaining the application of cervical traction for 1 min, slider neural mobilizations was applied during traction to mobilize and restore the normal structure and function of the nerve root that causing radiculopathy. Even though there are no standard parameters have been reported for cervical traction, it is speculated that traction causes distraction of articular surfaces, unloads the components of the spine by stretching muscles, ligaments, reduce adhesions within the dural sleeve, relieve nerve root compression within the central foramina, decreases pressure within intervertebral discs, relieves tonic muscle contraction and improves vascular status within the epidural space and perineural structures. Neural tissue mobilization techniques focus on restoring the ability of the nervous system to tolerate the normal compressive, friction, and tensile forces associated with daily and sport activities. The techniques used in this study were tensioning technique, Sliding technique and Single Joint Movement technique which are commonly

used Neural Mobilization techniq-ues. Sliding techniques during traction allows large range neurally non-aggressive movements. The clinical assumption is that these sliding techniques result in a larger longitudinal excursion of the nerve with a minimal increase in strain on impinged or tensed nerve. A gliding/tensioning technique may reduce intraneural swelling and circulatory compromise viafluctuating effects on intraneural pressure. Nerve gliding is induced by elongation of the nerve bed which elongates the nerve, increases the nerve tension and intraneural pressure reducing the intraneural blood flow in the oedematous neuropathies. Dynamically altering intraneural pressure may result in a 'pumping action' or 'milking effect' with beneficial effects on nerve hydration as it facilitates evacuation of the intra neural oedema when correctly applied and hence brings about a reduction in symptoms. The effects of these techniques individually have been explored in many RCTs and systematic reviews. In a single case study by Christos Savva et.al found that cervical traction combined with neural mobilization significantly shown effective in improving pain and disability in a patient with cervical radiculopathy. In Group B (Traction Group), there were significant improvement in post intervention measurements and this could be because of Mechanical cervical traction received for 12 sessions that might have shown decrease in pain level and perceived disability in patients with cervical radiculopathy. In Group C (Neural Mobilization Group) the improvement could be because of Neural Mobilization Techniques that used to normalize the CNRs (cervical nerve root) structure and function via the possible reduction of nerve adherence, facilitation of nerve gliding and decreased neural mechanosensitivity. When the NPRS score and NDI means were compared at pre-intervention and 2<sup>nd</sup> week post intervention between the groups there was no statistically significant difference between the groups. When the means of 4<sup>th</sup> week postintervention were compared between three group there was a statistically significant difference between the groups. There is significant improvement in severity of radicular Symptoms measured by Global Rating of Change Scale between the groups. Group A subjects shown greater percentage of reduction in severity of radicular Symptoms than Group B and C. These improvements were obtained without use of any conventional treatment techniques like exercises and modalities. The improvements obtained in Group A subjects were -71.33% in NPRS and - 59.71% in NDI and with greater percentage of reduction in radicular symptoms following 12 sessions of treatment during 4 weeks of duration. Therefore, based on findings it was found that the simultaneous application of mechanical cervical traction along with neural mobilization receivedby Group A subjects found to be more effective than the Groups B and C subjects. Therefore, the present study rejects null hypothesis.

Limitations of the Study: This study sample selection was considered age group between 45 to 55 only, thus results were not generalize the whole population. Dosage of treatment parameters of the combined treatment techniques in the study was not standardized according to individual patients. Effectiveness wasbased on subjective outcome measures therefore objective outcome measures were not used. The long term effect was not found.

## CONCLUSION

Simultaneous application of mechanical cervical traction with neural mobilization is more effective in improvingpain, functional disability and severity of radicular symptoms than mechanical cervical traction and neural mobilization alone for subjects with Unilateral Cervical Radiculopathy. The simultaneous application of mechanical cervical traction along with neural mobilization is recommended in the treatment of Unilateral Cervical Radiculopathy.

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