

INTRODUCTION

- Pain is a primary reason for seeking care from physical therapists.¹
- Point prevalence of chronic widespread pain (CWP): 10.6% 11.8%.²
- Neurodynamic mobilization (NDM) is often used clinically as an effective intervention for pain³⁻⁴ – with both local and widespread hypoalgesic mechanisms having been proposed. However, there is limited and conflicting research supporting these mechanisms.
- Efficacy and effectiveness of NDM for those with CWP warrants further investigation, but further evidence for efficacy in asymptomatic populations needs to be established.

AIM and HYPOTHESES

Aim: To assess the immediate local and widespread hypoalgesic effects of NDM applied to the upper extremity of asymptomatic subjects.

- Subjects receiving NDM will exhibit greater changes in local and widespread QST measures compared to those receiving sham NDM.
- Subjects receiving sliders will exhibit greater changes in local and widespread QST measures compared to those receiving tensioners.

METHODS

- Double-blind randomized controlled trial (Brenau IRB: 1208684-5)
- 60 asymptomatic subjects ages 18-65 randomized to 1 of 3 groups:
- Slider (n=20): 2 bouts of 10 "slider" NDMs targeting the left (L) median nerve in the ULNT2 position as described by Butler³
- Tensioner (n=20): 2 bouts of 10 "tensioner" NDMs targeting the L median nerve in the ULNT2 position as described by Butler³
- Sham (n=20): 2 bouts of 10 sham NDMs as described previously by Beneciuk et al⁵
- **Baseline quantitative sensory testing (QST) was conducted** bilaterally at the points depicted in Figure 1 immediately preintervention by an examiner blinded to group allocation.
- QST measures included measures of pressure pain threshold (PPT), thermal pain threshold (TPT), and thermal pain tolerance (TPTol).
- Subjects received the allocated intervention immediately followed by post-intervention testing by the same blinded examiner who conducted the baseline testing.
- Local QST measures reflect measures from the L thenar eminence (median nerve sensory field), while Widespread QST measures were calculated using the mean of the remaining 5 testing points.
- Data Analysis: Following assessment of normality via Shapiro-Wilk tests, within-group change was assessed using paired t-tests or Wilcoxon Signed Rank tests; and between-group differences in change were assessed using Kruskal-Wallis H-tests.

- 3. Butler DS. *Neurodynamic Techniques*. Adelaide, Australia: NOI Group 2005. NOI Group Publications; 2005.

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A) Thenar Eminence **B)** Dorsal aspect of 1st CMC

	<u>Pre</u> (mean ± SD)	<u>Post</u> (mean ± SD)	<u>Pre-Post Change</u> (Mean, 95% conf. interval)	<u>P Value for Within-</u> <u>Group Change</u>	P Value for Between-Group Difference
LocalPPT (kgf) - Slider ⁺ - Tensioner ⁺⁺ - Sham ⁺	5.80 ± 1.93 5.87 ± 2.54 4.79 ± 2.00	5.50 ± 2.25 5.50 ± 2.32 4.52 ± 1.77	-0.30 (-1.08 to 0.47) -0.36 (-1.36 to 0.64) -0.27 (-0.86 to 0.32)	.43 .85 .36	.67
WidespreadPPT (kgf) - Slider + - Tensioner + - Sham +	5.97 ± 2.02 6.74 ± 2.43 5.08 ± 1.35	6.75 ± 2.42 6.85 ± 2.37 5.39 ± 2.08	0.78 (0.14 to 1.42) 0.10 (-0.38 to 0.58) 0.31 (-0.32 to 0.95)	.02* .66 .32	.30
LocalTPT (°C) - Slider ⁺ - Tensioner ⁺ - Sham ⁺	41.76 ± 3.23 42.27 ± 3.48 42.00 ± 3.45	43.36 ± 3.59 44.28 ± 3.02 42.56 ± 2.93	1.60 (0.50 to 2.70) 2.01 (0.71 to 3.30) 0.56 (-0.71 to 1.83)	.01* .004* .37	.11
WidespreadTPT (°C) - Slider + - Tensioner + - Sham +	42.51 ± 2.88 43.00 ± 2.92 41.66 ± 2.94	43.94 ± 2.78 44.34 ± 2.88 43.07 ± 2.57	1.43 (0.73 to 2.14) 1.33 (0.64 to 2.02) 1.41 (0.62 to 2.20)	<.001* .001* .001*	.88
LocalTPTol (°C) - Slider ⁺⁺ - Tensioner ⁺⁺ - Sham ⁺	46.97 ± 3.23 47.91 ± 2.06 47.46 ± 2.15	48.15 ± 2.01 48.86 ± 1.50 48.07 ± 2.04	1.18 (0.49 to 1.87) 0.95 (0.20 to 1.69) 0.61 (0.13 to 1.08)	.001* .004* .02*	.51
WidespreadTPTol (°C) - Slider + - Tensioner ++ - Sham + Within-group analysis via +paired a	47.59 ± 1.90 47.79 ± 1.77 47.06 ± 2.00 t-test or <i>++Wilcoxon signed ra</i>	48.45 ± 1.45 48.44 ± 1.63 47.67 ± 1.68 nk test; Between-group a	0.86 (0.56 to 1.16) 0.65 (0.31 to 0.98) 0.61 (0.28 to 0.94) Inalysis via Kruskal-Wallis H-test	<.001* .001* .001*	.34

Reference List

1. Fruth SJ. Fundamentals of the Physical Therapy Examination: Patient Interview and Tests & Measures. Burlington, MA: Jones & Bartlett Learning; 2013. 2. Mansfield KE, Sim J, Jordan JL, Jordan KP. A systematic review and meta-analysis of the prevalence of chronic widespread pain in the general population. Pain. 2016;157(1):55-64. doi:10.1097/j.pain.000000000000014

4. Ellis RF, Hing WA. Neural mobilization: a systematic review of randomized controlled trials with an analysis of therapeutic efficacy. J Man Manip Ther. 2008;16(1):8-22. doi:10.1179/106698108790818594 5. Beneciuk JM, Bishop MD, George SZ. Effects of upper extremity neural mobilization on thermal pain sensitivity: a sham-controlled study in asymptomatic participants. J Orthop Sports Phys Ther. 2009;39(6):428-438. doi:10.2519/jospt.2009.2954

FIGURE 1. QST testing points (tested bilaterally)



CONCLUSION / DISCUSSION

- than sham NDM in producing a hypoalgesic effect.
- does result in some level of neurodynamic mobilization.
- local, regional, and widespread pain syndromes.

RESULTS



• Although statistically significant within-group changes were observed for multiple QST variables, there was no significant interaction between groups (Table 1).

• These findings suggest that in asymptomatic subjects, NDM may be no more effective

• However, one could argue that the sham NDM utilized in this study and elsewhere⁵

• Future studies should explore alternate methods of sham NDM and should assess for hypoalgesic effects of NDM in various symptomatic populations – such as those with