

RESEARCH REPOSITORY

This is the author's final version of the work, as accepted for publication following peer review but without the publisher's layout or pagination. The definitive version is available at:

http://dx.doi.org/10.1016/j.physio.2016.05.002

Koppenhaver, S.L., Walker, M.J., Rettig, C., Davis, J., Nelson, C., Su, J., Fernandez-de-las-Penas, C. and Hebert, J.J. (2017) The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study. Physiotherapy, 103 (2). pp. 131-137.

http://researchrepository.murdoch.edu.au/id/eprint/34840/



Copyright © 2017 Elsevier Ltd.

Accepted Manuscript

Title: The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study

Author: Shane L. Koppenhaver Michael J. Walker Charles Rettig Joel Davis Chenae Nelson Jonathan Su Cesar Fernández-de-las-Peñas Jeffrey J. Hebert



PII:	S0031-9406(16)30022-0
DOI:	http://dx.doi.org/doi:10.1016/j.physio.2016.05.002
Reference:	PHYST 907
To appear in:	Physiotherapy
Received date:	17-7-2015
Accepted date:	13-5-2016

Please cite this article as: Koppenhaver SL, Walker MJ, Rettig C, Davis J, Nelson C, Su J, Fernández-de-las-Peñas C, Hebert JJ, The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study, *Physiotherapy* (2016), http://dx.doi.org/10.1016/j.physio.2016.05.002

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

1	
2	
3 4 5 6	The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study
7	Shane L. Koppenhaver, PhD, PT ¹
8	Michael J. Walker, DSc, PT ²
9	Charles Rettig, DPT ³
10	Joel Davis, DPT ³
11	Chenae Nelson, DPT ³
12	Jonathan Su, DPT ³
13	Cesar Fernández-de-las-Peñas PT, PhD ⁴
14	Jeffrey J. Hebert, PhD ⁵
15	
16	¹ Associate Professor, U.S. Army-Baylor University Doctoral Program in Physical
17	Therapy, San Antonio, Texas; ² Associate Professor, South College Doctor of Physical
18	Therapy Program, Knoxville, Tennessee; ³ Physical Therapy Student during study,
19	U.S. Army-Baylor University Doctoral Program in Physical Therapy, San Antonio,
20	Texas; ⁴ Department of Physical Therapy, Occupational Therapy, Physical Medicine
21	and Rehabilitation, Universidad Rey Juan Carlos (URJC), Alcorcón, Madrid, Spain;
22	⁵ Associate Dean (Research), School of Psychology and Exercise Science, Murdoch
23	University, Perth, Australia.

25	This study was	performed at t	he Center for	Physical Ther	apy Research at th	e U.S. Army-
	2	1		2		J

- 26 Baylor University Doctoral Program in Physical Therapy, San Antonio, Texas and was
- approved by the Institutional Review Board of Brooke Army Medical Center.

28

- 29 The views expressed herein are those of the authors and do not reflect the official policy
- 30 or position of BAMC, the U.S. Army Medical Department, the U.S. Army Office of the
- 31 Surgeon General, the Department of the Army, Department of the Air Force, Department
- 32 of Defense, or the U.S. Government.
- 33
- 34 Corresponding Author:
- 35 Shane Koppenhaver
- 36 <u>shanekoppenhaver@mac.com</u>
- 37 210-722-3671
- 38
- 39 *Ethical Approval: Brooke Army Medical Center IRB

40

The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study

44

45 WORDS: 266

46 Abstract

47 **Objective:** To investigate the relationship between dry needling-induced twitch response 48 and change in pain, disability, nociceptive sensitivity, and lumbar multifidus muscle **49** function, in patients with low back pain (LBP). **Design:** Quasi-experimental study. 50 Setting: Department of Defense academic institution. Participants: Sixty-six patients 51 with mechanical LBP (38 men, 28 women, age: 41.3 [9.2] years). Interventions: Dry 52 needling treatment to the lumbar multifidus muscles between L3-L5 bilaterally. Main 53 Outcome Measures: Examination procedures included numeric pain rating, the 54 Modified Oswestry Disability Index, pressure algometry, and real-time ultrasound 55 imaging assessment of lumbar multifidus muscle function before and after dry needling 56 treatment. Pain pressure threshold (PPT) was used to measure nocioceptive sensitivity. 57 The percent change in muscle thickness from rest to contraction was calculated to 58 represent muscle function. Participants were dichotomized and compared based on 59 whether or not they experienced at least one twitch response on the most painful side and 60 spinal level during dry needling. **Results:** Participants experiencing local twitch response 61 during dry needling exhibited greater immediate improvement in lumbar multifidus 62 muscle function than participants who did not experience a twitch (thickness change with 63 twitch: 12.4 [5.7]%, thickness change without twitch: 5.7 [10.5]%, mean difference 64 adjusted for baseline value, 95%CI: 4.4 [1.2, 7.5]%). However, this difference was not

present after 1-week, and there were no between-groups differences in disability, pain
intensity, or nociceptive sensitivity. Conclusions: The twitch response during dry
needling might be clinically relevant, but should not be considered necessary for
successful treatment.
Key Words: Dry needling; low back pain; paraspinal muscles; muscle contraction;
ultrasonography.
Abbreviations: Low back pain (LBP); Modified Oswestry Disability Index (ODI)

The association between dry needling-induced twitch response and change in pain and muscle function in patients with low back pain: A quasi-experimental study

80

81 INTRODUCTION

Dry needling is a therapeutic procedure comprising of the insertion of a thin filiform needle directly into myofascial trigger points [1]. Clinical trials examining the effectiveness of dry needling have reported immediate and short-term pain relief and functional improvement for a wide range of musculoskeletal conditions [2–7]. Yet, recent systematic reviews have concluded that evidence for dry needling effectiveness is limited, owing to poor methodological quality and clinical heterogeneity among included trials [8–13].

89 Potentially important sources of clinical heterogeneity involve the differences in 90 dry needling technique including the role of the local twitch response [14]. A twitch response occurs when there is a brisk, involuntary contraction within the muscle being 91 92 needled [15]. It is believed that the twitch response results from a spinal reflex, following 93 the mechanical stimulation introduced by the needle [16,17]. Studies have demonstrated 94 both electrical and biochemical changes after eliciting twitch responses [14,18]. The 95 twitch response is often used to confirm the presence of trigger points which frequently 96 drives both patient selection and treatment parameters [19]. Likewise, many practitioners 97 assume that the elicitation of a twitch response during dry needling represents evidence 98 of trigger point "inactivation" and is necessary for achieving a successful clinical 99 outcome. However, few studies have examined the potential relationship between dry 100 needling-induced local twitch response and clinical improvements [5,16]. Moreover, the 101 results of these studies conflict, with one reporting immediate changes in pain and range

of motion only in participants experiencing twitch response [16] and the other reporting
no differences in quality of life based on local twitch response and only differences in
pain after 4 weeks [5]. Additionally, both of these studies exhibited important limitations
such as procedures that were not standardized [16] and small sample sizes [5].

The lumbar multifidus muscle has been shown to play an important role for normal function of the lumbar spine and has been implicated clinically in patients with low back pain (LBP) [20,21]. No prior studies have examined the effect of twitch response during dry needling on lumbar multifidus muscle function and clinical outcome in patients with LBP. Therefore, the purpose of this study was to explore the relationship between dry needling-induced local twitch response and change in pain, LBP-related disability, nociceptive sensitivity, and lumbar multifidus muscle function in patients with LBP.

113

114 METHODS

115 Study Design

116 This study was a pre-planned secondary analysis of data from a quazi-117 experimental study investigating changes in lumbar multifidus muscle function and 118 nociceptive sensitivity in LBP patient responders vs. non-responders after dry needling 119 treatment [22]. The study protocol was approved by the Institutional Review Board of 120 Brooke Army Medical Center and all participants provided written informed consent 121 prior to study enrollment. The study entailed two visits consisting of the same procedures 122 for all participants. Visit #1 included self-report questionnaires, baseline history and 123 physical examination, dry needling treatment to lumbar multifidus muscles, and pre- and 124 post-needling pain measures, pressure algometry and real-time ultrasound imaging

assessment of lumbar multifidus muscle function. Visit #2 occurred approximately one
week after visit #1 and included repeat self-report questionnaires, pressure algometry,

127 and real-time ultrasound imaging assessment of lumbar multifidus muscle function.

128 Study Participants

129 Study participants were recruited through print and email advertising within the 130 San Antonio Military Healthcare System. We recruited participants between the ages of 18 and 60 years, with current LBP (defined as pain located between the 12th rib and 131 132 buttocks), and a minimum Modified Oswestry Disability Index (ODI) score of at least 133 20/100. Potential participants were excluded if they were pregnant, taking anticoagulant 134 medication, or displayed signs of lumbar radiculopathy or non-musculoskeletal pathology 135 (e.g. cancer, infection). Additionally, we excluded individuals who reported a history of 136 lumbar spine surgery, bleeding disorder, and those who had performed trunk stabilization 137 exercises or received manual therapy to the lumbar region in the preceding month. All 138 individuals provided written informed consent prior to study enrollment.

139 Procedures

140 All participants underwent a standardized history and physical examination based 141 on the tests and measures associated with the treatment-based classification system [23]. 142 During the examination, participants nominated the most painful side of their low back 143 region (right or left). If the participant's sides were equally painful, then the symptomatic 144 side was chosen at random. Pain intensity and pain-related disability were self-reported 145 by each participant. The ODI consists of scores ranging from 0 to 100, with higher scores 146 representing higher levels of disability, and has previously been found to be both reliable 147 and responsive to change [24,25]. An 11-point numeric pain rating scale was used to

quantify participants' current back pain intensity. The numeric pain rating scale has been
shown to be reliable and responsive (minimally important difference = 2 points) in
patients with LBP [26,27].

151 *Pressure Algometry*

152 Pressure algometry was used to determine the most painful spinal level at baseline 153 and as a measure of nociceptive sensitivity identified by the pain pressure threshold 154 (PPT). PPT is the minimal amount of pressure that produces pain [28] and is used to 155 assess abnormalities in nociceptive processing or hyperalgesia [28,29]. A digital pressure 156 algometer (Wagner Force Ten FDX, Wagner Instruments, Greenwich, CT) was used to 157 measure PPT at L3, L4, and L5 paraspinal muscles on the most symptomatic side. An 158 examiner applied the pressure algometer perpendicular to the muscle belly of lumbar 159 multifidus, approximately 1.5 cm lateral to the spinous process. The algometer was 160 advanced at a rate of approximately 5N/s and participants were instructed to verbally 161 signal when they first perceived the force change from "pressure" to "pain." Previous 162 studies have found PPT measures to be highly reliable and responsive to change [30,31]. 163 PPT at each location was taken three times and averaged to reduce measurement error.

164 Ultrasound Imaging Assessment of Muscle Function

Real-time ultrasound imaging measures muscle function by quantifying the change in muscle thickness from resting to contracted states [32,33]. Studies have found ultrasound measurements of the lumbar multifidus musculature to be reliable (minimal detectable change = 1.6mm to 2.8mm) [33] and valid [34]. Images of the lumbar multifidus muscle were acquired at rest and during a sub-maximal contraction at levels L4/5 and L5/S1 on the more symptomatic side following techniques outlined in previous

171 work [33,35]. All ultrasound images were obtained using a Sonosite Titan (Sonosite Inc. 172 Bothell, WA) with a 60mm 5MHz curvilinear array by a trained examiner that was 173 blinded as to whether a participant experienced a twitch or not during dry needling. A 174 contralateral arm lift maneuver while holding a hand weight normalized to body mass 175 was used to elicit a 30% maximal voluntary isometric contraction [32]. One practice lift 176 was performed followed by 3 image acquisitions at rest and during the contralateral arm 177 lift. Images were exported and measured offline using Image J software (Wayne 178 Rasband, National Institutes of Health, USA). Muscle thickness was measured as the 179 distance between the posterior-most portion of the L4/L5 or L5/S1 facet joint and the 180 fascial plane between the muscle and subcutaneous tissue. By using Image J's automatic measurement function, the examiner was additionally blinded to thickness values during 181 182 measurement. The 3 measures of each condition (rest and contraction) were averaged to

183 reduce measurement error [36].

184 Dry Needling Treatment

All participants underwent a single session of dry needling therapy performed by one of two experienced physical therapists who were fellowship trained in orthopedic manual therapy, trained in dry needling, and blinded to baseline assessment outcomes. The examiner palpated the lumbar multifidus muscles to identify the presence of trigger points, which we defined as a palpable and painful nodules in the muscle tissue [37].

190 The needling technique included insertion of a sterile, disposable, 0.30x50 mm or 191 0.30x60 mm solid filament needle (Seirin Corp., Shizuoka, Japan) into the lumbar 192 multifidus muscles at the L3, L4 and L5 spinal levels bilaterally (Figure 1). Needles were 193 inserted approximately 1.5 cm lateral to the spinous process at each segmental level in a

194 posterior to anterior direction. After piercing the skin, the needles were directed into the 195 lumbar multifidus muscle with a slight inferior-medial angle (approximately 20 degrees) 196 to the depth of the lumbar lamina and further localized towards trigger points when 197 detected. Each segment was treated once on each side, with needle insertion lasting 198 approximately 5-10 seconds. "Sparrow pecking" (in and out motion) and "coning" (small 199 redirections of needle angle) techniques were utilized in an attempt to elicit as many 200 twitch responses as possible [38]. The presence of local twitch response was considered 201 to occur if at least one visible or palpable muscle twitch was observed by the examiner or 202 reported by the participant.

203 Statistical analysis

204 The most symptomatic side (right vs. left) was established during the baseline 205 assessment. To further localize analysis to the most painful area, the most symptomatic 206 level on the more symptomatic side was identified by the spinal level (L4 vs. L5) with the 207 lowest PPT for each participant. Participants were then categorized based on whether or 208 not local twitch response was elicited on the most symptomatic side and spinal level. 209 Baseline characteristics were compared with independent t-tests for normally distributed 210 continuous-level variables, Man-Whitney U test for non-normally distributed continuous-211 level variables, and Chi-square tests for categorical variables.

Muscle function was calculated for the most painful spinal level (L4/5 vs. L5/S1) at each time point (baseline, immediately after needling, and 1 week after needling) using the equation [contracting thickness-resting thickness]/resting thickness. PPT was averaged across spinal levels (L3, L4, L5) at each time point to represent the dependent variable of nociceptive sensitivity. Separate analysis of covariance models were used to

examine for differences in each dependent variable (ODI, pain, PPT, muscle activation)
at each time point (immediately after needling, 1 week after needling) after adjusting for
baseline values. All data were analyzed with IBM SPSS Version 21 software (Chicago,
IL) using a pre-specified alpha of 0.05.

221

222 RESULTS

Two hundred and sixty individuals were screened for study inclusion. One 223 224 hundred and eighty eight were excluded, most commonly for having an ODI score of less 225 than 20%. Of the 72 participants enrolled in the study, 6 individuals failed to return for 226 the follow up visit, leaving complete data on 66 participants. The complete participant 227 flow chart has been published elsewhere [22]. Of the 66 participants, 61 (92%) exhibited 228 at least one twitch response (and usually more than one) during treatment. Thirty-five 229 participants (53%) experienced at least one twitch at the most symptomatic side and 230 spinal level during dry needling. Follow-up reassessment occurred a mean of 6.3 (SD: 231 1.9) days after the dry needling. Baseline demographic and clinical history information, 232 stratified by twitch response status is displayed in Table 1. There were no baseline 233 differences between participants that exhibited local twitch response and those that did 234 not at baseline.

Participants experiencing local twitch response demonstrated greater immediate improvement in lumbar multifidus muscle function than those who did not experience a twitch. However, this difference was not present after 1-week (**Table 2, Figure 2**). There were no between-groups differences in disability, pain intensity, or nociceptive sensitivity (**Table 2, Figure 3**).

240

241 **DISCUSSION**

242 Although clinicians often view the elicitation of local twitch response during dry 243 needling as a primary goal and indicator of successful treatment there is scarce evidence 244 supporting this assertion [16,39]. Therefore, the purpose of the current study was to 245 explore the relationship between dry needling-induced twitch response and changes in 246 pain, LBP-related disability, nociceptive sensitivity, and lumbar multifidus muscle 247 function in patients with LBP. Our primary finding was that twitch response elicited on 248 the most painful side and spinal level during dry needling is related to an immediately 249 improvement in lumbar multifidus activation, but not pain, nociceptive sensitivity, LBP-250 related disability, or lasting improvements in muscle function.

251 Few other studies have investigated the clinical relevance of the local twitch 252 response. The earliest study by Hong et al. [16] was focused on comparing the effect of 253 dry needling vs. lidocaine injection to the upper trapezius muscle on pain, PPT, and 254 cervical range of motion in 58 patients with myofascial pain syndrome. A secondary 255 analysis compared outcomes in those that experienced twitch response (n=41) to those 256 that did not (n=17). Somewhat contradictory to that of the current study, Hong et al. [16] 257 found statistically significant changes in pain, PPT, and range of motion in participants 258 that experienced a local twitch response and little to no statistically significant changes in 259 participants that did not experience a twitch response immediately after dry needling. 260 However, this study had methodological limitations, such as lack of blinding, not 261 standardizing procedures, and they did statistically compare the responses in those that 262 experienced a twitch response to those that did not.

263 A more recent study by Tekin et al. [5] compared changes in pain and quality of 264 life after dry needling or sham dry needling to the upper back in 39 subjects with 265 myofascial pain syndrome. A secondary analysis of the trial was performed in the 22 266 subjects that received dry needling to compare outcomes in those that experienced twitch 267 response (n = 9) to those that did not (n=13). Although they did not find any difference in 268 quality of life (SF-36), subjects that experienced local twitch during dry needling 269 demonstrated larger improvements in pain at 4 weeks, but not after 1 week. Further, this 270 difference at 4 weeks was of sufficient magnitude to be considered clinically significant 271 (approximately 2 points on VAS).

272 In the last and only study to include muscles of the low back region, Rha et al. 273 [39] evaluated the ability of ultrasound imaging to detect twitch responses during trigger 274 point injection to upper trapezius, erector spinae, or quadratus lumborum muscles in 41 275 patients with myofascial pain syndrome. A secondary analysis within their primary study 276 found a statistically larger immediate reduction in pain in those participants that exhibited 277 local twitch response than those that did not during the injection. Similar to Tekin et al. 278 [5], the magnitude of difference in pain reduction was sufficiently large enough to be 279 considered clinically significant (2.6 to 2.9 points on the VAS).

Of note, there appears to be large variability in prevalence rates of a local twitch response between the previously discussed studies [5,16] and the current study. While the majority of participants experienced a twitch response in the current study (92.4% overall and 53.0% at the most symptomatic side and spinal level) and the earlier one by Hong et al. [16] (71%), Tekin et al. [5] reported twitch responses in only a minority of subjects (41%). Although the reason for this difference is unknown, it may be at least partially due

to the muscle or region being treated with dry needling (low back vs. upper back andtrapezius).

288 When the findings of the current study are added to the findings of these few prior studies [5,16,39], it appears that local twitch response during needling may be related to 289 290 an immediate improvement in muscle function and may or may not be related to 291 clinically important reductions in pain after dry needling. However, twitch response is 292 unlikely to be related to changes in pain-related disability or quality of life. This suggests 293 that twitch response during dry needling might be clinically relevant, but that it should 294 not be considered a "hallmark" sign of dry needling or "necessary" for successful 295 treatment.

296 The primary limitation of the current study concerns the inherent challenges of 297 identifying local twitch response, especially in the lumbar multifidus muscle. Inter-rater 298 reliability of twitch response identification has been reported to be low (kappa = -0.02 to 299 0.18) regardless of the muscle examined or the level of training of the examiner [40]. 300 When comparing the detection of twitch responses via visual inspection to 301 ultrasonography, Rha et al. [39] found that visual inspection was able to detect all twitch 302 responses in the upper trapezius muscle, and most, but not all of the local twitch 303 responses in the lower back musculature (erector spinae and quadratus lumborum) when 304 compared to ultrasonography. Future research should evaluate the clinical relevance of 305 twitch response using more superficial muscles (e.g. infraspinatus) and/or using more 306 accurate identification measures (e.g. ultrasonography or EMG).

307 Other salient limitations of the current study were the lack of our ability to blind the 308 participants and the relatively short reassessment period (1 week). In the author's

309 experience, a local twitch response is a fairly intense sensation to patients that is often 310 described as similar to a "jolt of lighting." Therefore, it is possible that participants 311 experienced a placebo effect from the twitch response. We attempted to minimize this 312 effect by having all outcomes obtained by examiners that were blinded to whether or not 313 participants experienced twitch response. Moreover, the only outcome that showed a 314 difference based on local twitch response was lumbar multifidus muscle function, which 315 arguably would be the least likely measure affected by placebo. Lastly, it is possible that 316 the local twitch response was related to longer term (> 1 week) changes in pain and/or 317 disability as reported by Tekin et al. [5] as we only reassessed participants 1 week after 318 dry needling. However, considering that altered muscle function was only associated with 319 the twitch response immediately after, and not 1 week after, dry needing, this is not likely 320 the case. Alternatively, it could be that dry needling treatment would have more lasting 321 effects when followed by some additional muscle activation or strengthening exercises.

322

323 CONCLUSION

Local twitch response elicited on the most painful side and spinal level during dry needling appears to be related to immediately improve lumbar multifidus function, but not pain, nociceptive sensitivity, LBP-related disability, or lasting improvements in muscle function. This suggests that the local twitch response during dry needling might be clinically relevant, but that it should not be considered as a "hallmark" sign of dry needling or "necessary" for successful treatment.

330

332	*Funding: None.
-----	-----------------

333 *Conflict of Interest: None

334

335 **REFERENCES**

- 336 Kalichman L, Vulfsons S. Dry needling in the management of musculoskeletal pain. [1] 337 J Am Board Fam Med 2010;23:640-6. doi:10.3122/jabfm.2010.05.090296. 338 [2] Dıraçoğlu D, Vural M, Karan A, Aksoy C. Effectiveness of dry needling for the 339 treatment of temporomandibular myofascial pain: a double-blind, randomized, 340 placebo controlled study. J Back Musculoskelet Rehabil 2012;25:285-90. 341 doi:10.3233/BMR-2012-0338. 342 [3] Llamas-Ramos R, Pecos-Martín D, Gallego-Izquierdo T, Llamas-Ramos I, Plaza-343 Manzano G, Ortega-Santiago R, et al. Comparison of the Short-Term Outcomes 344 Between Trigger Point Dry Needling and Trigger Point Manual Therapy for the 345 Management of Chronic Mechanical Neck Pain: A Randomized Clinical Trial. 346 Journal of Orthopaedic & Sports Physical Therapy 2014;44:852-61. 347 doi:10.2519/jospt.2014.5229. 348 [4] Mejuto-Vázquez MJ, Salom-Moreno J, Ortega-Santiago R, Truyols-Domínguez S, 349 Fernández-de-las-Peñas C. Short-Term Changes in Neck Pain, Widespread Pressure 350 Pain Sensitivity, and Cervical Range of Motion After the Application of Trigger Point Dry Needling in Patients With Acute Mechanical Neck Pain: A Randomized 351 352 Clinical Trial. Journal of Orthopaedic & Sports Physical Therapy 2014;44:252–60. 353 doi:10.2519/jospt.2014.5108. 354 Tekin L, Akarsu S, Durmus O, Cakar E, Dincer Ü, Kıralp MZ. The effect of dry [5] 355 needling in the treatment of myofascial pain syndrome: a randomized double-356 blinded placebo-controlled trial. Clinical Rheumatology 2012;32:309-15. 357 doi:10.1007/s10067-012-2112-3. 358 Itoh K, Saito S, Sahara S, Naitoh Y, Imai K, Kitakoji H. Randomized Trial of [6] 359 Trigger Point Acupuncture Treatment for Chronic Shoulder Pain: A Preliminary 360 Study. Journal of Acupuncture and Meridian Studies 2014;7:59-64. 361 doi:10.1016/j.jams.2013.02.002. 362 Chou L, Hsieh Y, Chen H, Hong C, Kao M, Han T. Remote Therapeutic [7] 363 Effectiveness of Acupuncture in Treating Myofascial Trigger Point of the Upper Trapezius Muscle. Am J Phys Med Rehabil 2011. 364 365 doi:10.1097/PHM.0b013e3182328875.
- Kietrys DM, Palombaro KM, Azzaretto E, Hubler R, Schaller B, Schlussel JM, et al.
 Effectiveness of Dry Needling for Upper-Quarter Myofascial Pain: A Systematic
 Review and Meta-analysis. Journal of Orthopaedic & Sports Physical Therapy
 2013;43:620–34. doi:10.2519/jospt.2013.4668.
- Tough E, White A. Effectiveness of acupuncture/dry needling for myofascial trigger
 point pain. Physical Therapy Reviews 2011;16:147–54.
- [10] Liu L, Huang Q-M, Liu Q-G, Ye G, Bo C-Z, Chen M-J, et al. Effectiveness of Dry
 Needling for Myofascial Trigger Points Associated With Neck and Shoulder Pain:

374		A Systematic Review and Meta-Analysis. Archives of Physical Medicine and
375		Rehabilitation 2015;96:944–55. doi:10.1016/j.apmr.2014.12.015.
376	[11]	Boyles R, Fowler R, Ramsey D, Burrows E. Effectiveness of trigger point dry
377		needling for multiple body regions: a systematic review. Journal of Manual &
378		Manipulative Therapy 2015:2042618615Y.0000000014.
379		doi:10.1179/2042618615Y.0000000014.
380	[12]	Ong J, Claydon LS. The effect of dry needling for myofascial trigger points in the
381		neck and shoulders: A systematic review and meta-analysis. Journal of Bodywork
382		and Movement Therapies 2013. doi:10.1016/j.jbmt.2013.11.009.
383	[13]	France S, Bown J, Nowosilskyj M, Mott M, Rand S, Walters J. Evidence for the use
384		of dry needling and physiotherapy in the management of cervicogenic or tension-
385		type headache: A systematic review. Cephalalgia 2014:34:994–1003.
386		doi:10.1177/0333102414523847.
387	[14]	Hsieh Y-L, Yang S-A, Yang C-C, Chou L-W. Dry Needling at Myofascial Trigger
388		Spots of Rabbit Skeletal Muscles Modulates the Biochemicals Associated with Pain.
389		Inflammation, and Hypoxia, Evidence-Based Complementary and Alternative
390		Medicine 2012:2012:1–12. doi:10.1155/2012/342165.
391	[15]	Chou L-W, Kao M-J, Lin J-G. Probable Mechanisms of Needling Therapies for
392		Myofascial Pain Control. Evidence-Based Complementary and Alternative
393		Medicine 2012;2012:1–11. doi:10.1155/2012/705327.
394	[16]	Hong C. Lidocaine injection versus dry needling to myofascial trigger point. The
395		importance of the local twitch response. Am J Phys Med Rehabil 1994;73:256–63.
396	[17]	Rivner MH. The neurophysiology of myofascial pain syndrome. Curr Pain
397		Headache Rep 2001;5:432–40.
398	[18]	Hsieh Y-L, Chou L-W, Joe Y-S, Hong C-Z. Spinal cord mechanism involving the
399		remote effects of dry needling on the irritability of myofascial trigger spots in rabbit
400		skeletal muscle. Arch Phys Med Rehabil 2011;92:1098–105.
401		doi:10.1016/j.apmr.2010.11.018.
402	[19]	Huang Y-T, Lin S-Y, Neoh C-A, Wang K-Y, Jean Y-H, Shi H-Y. Dry needling for
403		myofascial pain: prognostic factors. J Altern Complement Med 2011;17:755–62.
404		doi:10.1089/acm.2010.0374.
405	[20]	Freeman M, Woodham M, Woodham A. The Role of the Lumbar Multifidus in
406		Chronic Low Back Pain: A Review. PM R 2010;2:142-6.
407		doi:10.1016/j.pmrj.2009.11.006.
408	[21]	MacDonald DA, Moseley GL, Hodges PW. The lumbar multifidus: does the
409		evidence support clinical beliefs? Man Ther 2006;11:254-63.
410	[22]	Koppenhaver SL, Walker MJ, Su J, McGowen JM, Umlauf L, Harris KD, et al.
411		Changes in lumbar multifidus muscle function and nociceptive sensitivity in low
412		back pain patient responders versus non-responders after dry needling treatment.
413		Manual Therapy 2015. doi:10.1016/j.math.2015.03.003.
414	[23]	Hebert JJ, Koppenhaver SL, Walker BF. Subgrouping Patients With Low Back
415		Pain. Sports Health: A Multidisciplinary Approach 2011;3:534–42.
416		doi:10.1177/1941738111415044.
417	[24]	Fritz JM, Irrgang JJ. A Comparison of a Modified Oswestry Disability
418		Questionnaire and the Quebec Back Pain Disability Scale. Phys Ther 2001;81:776–
419		88.

420	[25]	Ostelo R, Deyo R, Stratford P, Waddell G, Croft P, Von Korff M, et al. Interpreting
421		change scores for pain and functional status in low back pain: towards international
422		consensus regarding minimal important change. Spine 2008;33:90–4.
423		doi:10.1097/BRS.0b013e31815e3a10.
424	[26]	Childs JD. Piva SR. Fritz JM. Responsiveness of the numeric pain rating scale in
425		patients with low back pain. Spine 2005:30:1331–4.
426	[27]	Cleland JA, Whitman JM, Houser JL, Wainner RS, Childs JD, Psychometric
427	Γ.1	properties of selected tests in patients with lumbar spinal stenosis. The Spine
428		Journal 2012:12:921–31. doi:10.1016/i.spinee.2012.05.004.
429	[28]	Ylinen J. Pressure algometry. Aust J Physiother 2007:53:207.
430	[29]	Sterling M. Pressure algometry: what does it really tell us? J Orthop Sports Phys
431	Γ.1	Ther 2011:41:623–4. doi:10.2519/iospt.2011.0106.
432	[30]	Kinser A. Sands W. Stone M. Reliability and validity of a pressure algometer. J
433		Strength Cond Res 2009:23:312–4.
434	[31]	Walton DM, Macdermid JC, Nielson W, Teasell RW, Chiasson M, Brown L.
435		Reliability, standard error, and minimum detectable change of clinical pressure pain
436		threshold testing in people with and without acute neck pain. J Orthop Sports Phys
437		Ther 2011;41:644–50. doi:10.2519/jospt.2011.3666.
438	[32]	Kiesel K, Uhl T, Underwood F, Rodd D, Nitz A. Measurement of lumbar multifidus
439		muscle contraction with rehabilitative ultrasound imaging. Man Ther 2007;12:161–
440		6. doi:10.1016/j.math.2006.06.011.
441	[33]	Koppenhaver S, Hebert J, Fritz J, Parent E, Teyhen D, Magel J. Reliability of
442		rehabilitative ultrasound imaging of the transversus abdominis and lumbar
443		multifidus muscles. Arch Phys Med Rehabil 2009;90:87–94.
444		doi:10.1016/j.apmr.2008.06.022.
445	[34]	Koppenhaver S, Hebert J, Parent E, Fritz J. Rehabilitative ultrasound imaging is a
446		valid measure of trunk muscle size and activation during most isometric sub-
447		maximal contractions: a systematic review. Aust J Physiother 2009;55:153-69.
448	[35]	Koppenhaver S, Fritz J, Hebert J, Kawchuk G, Childs J, Parent E, et al. Association
449		between changes in abdominal and lumbar multifidus muscle thickness and clinical
450		improvement after spinal manipulation. J Orthop Sports Phys Ther 2011;41:389–99.
451		doi:10.2519/jospt.2011.3632.
452	[36]	Koppenhaver S, Parent E, Teyhen D, Hebert J, Fritz J. The effect of averaging
453		multiple trials on measurement error during ultrasound imaging of transversus
454		abdominis and lumbar multifidus muscles in individuals with low back pain. J
455		Orthop Sports Phys Ther 2009;39:604-11. doi:10.2519/jospt.2009.3088.
456	[37]	Dommerholt J. Dry needling ? peripheral and central considerations. J Man Manip
457		Ther 2011;19:223–7. doi:10.1179/106698111X13129729552065.
458	[38]	Itoh K, Katsumi Y, Hirota S, Kitakoji H. Effects of trigger point acupuncture on
459		chronic low back pain in elderly patientsa sham-controlled randomised trial.
460		Acupunct Med 2006;24:5–12.
461	[39]	Rha D, Shin JC, Kim Y-K, Jung JH, Kim YU, Lee SC. Detecting local twitch
462		responses of myofascial trigger points in the lower-back muscles using
463		ultrasonography. Arch Phys Med Rehabil 2011;92:1576–80.e1.

464 doi:10.1016/j.apmr.2011.05.005.

- 465 [40] Hsieh C-YJ, Hong C-Z, Adams AH, Platt KJ, Danielson CD, Hoehler FK, et al.
- 466 Interexaminer reliability of the palpation of trigger points in the trunk and lower 467 limb muscles. Archives of Deviced Medicine and Debabilitation 2000.81,258, (4)
- 467 limb muscles. Archives of Physical Medicine and Rehabilitation 2000;81:258–64.
- 468

Characteristics	Entire Sample	Twitch Response	No Twitch	P-value
	(n=66)	(n=35)	Response (n=31)	
Age (years)	41.3 (9.2)	40.6 (8.7)	42.2 (9.9)	0.480
Sex (% women)	42%	51%	32%	0.140†
BMI (kg/m ²)	28.8 (4.9)	28.1 (4.6)	29.2 (5.1)	0.294
ODI score (%)	31.2 (11.4)	31.9 (11.7)	30.4 (11.3)	0.598
Numeric pain rating scale for back (0-10)	4.7 (1.7)	4.9 (1.8)	4.5 (2.1)	0.399
Duration of symptoms (months)	9.2 (0.4, 98.9)*	6.4 (0.2, 135.3)*	9.7 (0.9, 209.5)*	0.699††

468 TABLE 1: Baseline Demographic and History Information

469

470 Abbreviations: BMI, body mass index; ODI, Modified Oswestry Disability Index.

471 Values are mean (SD) unless otherwise indicated.

Ċ

472 [†]P-value from a Chi-Square test.

473 * Median (interquartile range)

474 †† P-value from Man-Whitney U

475

- 476 **TABLE 2:** Immediate and 1-week changes in disability, pain, pain pressure threshold,
- 477 and lumbar multifidus muscle activation after dry needling
- 478

	Baseline	Immediately after dry	1-week after
		needling	dry needling
Oswestry Disability Question	naire (0-100)		•.0
Twitch Response	31.4 (11.4)		24.6 (12.6)
No Twitch Response	30.4 (11.3)		21.9 (14.3)
Adj. Mean Difference (95%	bCI)		1.3 (-4.2, 7.0);
			P=0.624
Numeric Pain Rating Scale (0	-10)		
Twitch Response	4.76 (1.69)	2.62 (1.74)	2.68 (2.01)
No Twitch Response	4.45 (2.13)	3.42 (2.63)	2.65 (2.03)
Adj. Mean Difference (95%	oCI)	-1.0 (-2.0, 0.0)	-0.1 (-1.0, 0.8)
		P=0.051	P=0.829
Pressure Pain Threshold (N/c	m ²)		
Twitch Response	6.32 (3.64)	6.97 (3.66)	6.96 (3.61)
No Twitch Response	6.59 (3.41)	7.10 (3.85)	7.60 (3.84)
Adj. Mean Difference (95%	oCI)	0.11 (-0.79, 1.02)	-0.53 (-1.83, 0.78)
		P=0.807	P=0.422
Muscle Activation (% thickne	ess change from re	st)	
Twitch Response	10.2 (9.8)	12.4 (5.7)	9.7 (11.0)
No Twitch Response	7.4 (13.6)	5.7 (10.5)	6.3 (8.4)
Adj. Mean Difference (95%	oCI)	4.4 (1.2, 7.5)	2.0 (-2.0, 6.0)
		P=0.007*	P=0.318

- 480 Adjusted Mean Differences are (Twitch Response No Twitch Response) adjusted based on
- 481 baseline values.
- 482 *Statistically significant at p < 0.01
- 483



- **FIGURE 1:** Dry needling technique to the lumbar multifidus muscle (using a simulated
- 486 needle for visibility).

- -





FIGURE 2: Percent activation of lumbar multifidus muscle during contra-lateral arm lift

analyzed by presence of local twitch response at the most symptomatic side and vertebral

- 503 level during dry needling.



