Trigger point dry needling: the data do not support broad applicability or robust effect

Kenneth Venere¹, Kyle Ridgeway²

Intermountain Homecare and Hospice, Salt Lake City, UT, USA, University of Colorado Hospital, Denver, CO, **USA**

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We read with great interest the recent systematic review by Boyles et al. entitled "Effectiveness of trigger point dry needling for multiple body regions: a systematic review".1 We admire the work done by the authors to conduct a systematic review on an intervention of growing popularity in our profession and appreciate their efforts to enhance understanding of this intervention. However, we strongly disagree with the authors' conclusion when they write: "The majority of high-quality studies included in this review show measured benefit from trigger point dry needling for MTrPs in multiple body areas, suggesting broad applicability of trigger point dry needling treatment for multiple muscle groups".

This strongly worded conclusion overstates the findings of the actual data and misleads casual readers into believing that the research supporting trigger point dry needling is quite robust. We contend that it is not. While there are some measured differences between groups in some studies, actual clinical benefits appear questionable, at best, and suggestions of applicability are currently unwarranted. The individual data of many trials included in the systematic review demonstrate one or a combination of the following limitations:

- (1) No statistically significant difference from sham treatments.2-6
- (2) Outcomes measuring immediate effects only.^{3,7-11}
- (3) Clinically irrelevant effects on pain and disability. 3,7,10-15
- (4) Large degrees of uncertainty due to wide confidence intervals. 12,16
- (5) Failure to control for significant confounding variables, such as natural history, regression to the mean and non-specific treatment effects. 4,5,7,8,10,13,17,18
- (6) Withdrawn from publication at the request of the editor.19
- (7) Required an erratum due to misreporting of kappa values.20

Specifically, only 47% of the included trials showed a statistically significant decrease in pain when compared to sham or alternative treatments, only 26% displayed a

Correspondence to: Kenneth Venere, Intermountain Homecare and Hospice. Email: Kenneth. Venere@imail.org

statistically significant decrease in disability and 42% did not include a sham or control intervention group. Of the 10 trials that did include a sham intervention, only three actually assessed the quality of blinding in the sham group. Further, 32% of the included trials investigated only the immediate effects (ranging from immediately post intervention to 72 h) of Trigger Point Dry Needling (TDN) which is a research design with remarkable limitations.²¹ Further, one of the nineteen studies was retracted at the request of the journal editor. These are significant methodological concerns that should give any clinician or researcher pause when interpreting this systematic review and/or considering the implementation of trigger point dry

The Cotchett et al. trial²² on trigger point dry needling for heel pain included in the present systematic review showcases several of the issues broadly prevalent in the dry needling literature. Boyles et al. state that "Cotchett et al.²² found significant improvements in pain and subjective foot health report for trigger point dry needling to the plantar foot as compared to sham needling". We feel this to be a misrepresentation of the Cotchett et al.²² trials' results. While there was a statistically significant difference found between dry needling and sham dry needling measured by the Foot Health Status Questionnaire, the results failed to achieve a previously determined minimally important difference. This makes the results' clinical relevance questionable. Also of note, the adjusted mean difference between dry needling and sham for first-step pain measured by a 100-mm visual analog scale barely met the minimally important difference and exhibited rather wide confidence intervals (mean -14.495%CI (-23.5 to -5.2), thus limiting the certainty of the results. Most importantly, the frequency of adverse events between the dry needling group and the sham group resulted in a number needed to harm one of three. That is, for every three patients treated with dry needling (as opposed to sham), one will develop an adverse event that he/she would not have experienced had the patient not been treated with dry needling. This is a troublesome finding when considering that the authors found that the number needed to treat to obtain a useful outcome to be four. With the above information in mind, qualifying the trial as finding "significant improvements in pain and subjective foot health" is dubious at best.

Beyond the trial data itself, those utilizing a myofascial trigger point approach to dry needling need to acknowledge the current issues with regards to a lack of consistent criteria defining a trigger point^{17,23} the questionable clinical importance, if any, of trigger points, and the poor reliability in trigger point identification.²⁴ Moreover, there are several foundational issues at the core of the trigger point construct including a lack of established validity and the tautological reasoning inherent in both trigger points and myofascial pain syndrome.^{25–27} These issues, while beyond the scope of this letter, cast significant doubt on the foundational premise of trigger points as a clinical meaningful entity and treatment target via dry needling or other means.

Given the data included in the systematic review it is premature, if not inaccurate, for the authors to conclude that dry needling has broad applicability in the treatment of pain. We urge the readers to take pause and assess not only the individual trials included, but the premise, plausibility and proposed mechanisms of trigger point dry needling. The above concerns regarding the data on trigger point dry needling suggest that robust conclusions of effectiveness and applicability are not currently possible.

We suggest that the authors amend their strongly worded conclusions to more closely match what the actual trial data suggest that despite strong anecdotal support²⁸ and positive reviews (that are flawed),^{1,29,30} the positive influence of dry needling is overstated in this systematic review.³¹

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Conflict of interest

None declared.

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