# TØRRNÅLING ("DRY NEEDLING")

Tough EA, White AR, Cummings TM, Richards SH, Campbell JL. Acupuncture and dry needling in the management of myofascial trigger point pain: A systematic review and meta-analysis of randomised controlled trials. Eur J Pain. 2009 Jan;13(1):3-10.

### Dry needling compared to placebo for myofascial trigger point pain

Patient or population: Myofascial trigger point pain

Intervention: Dry needling Comparison: Placebo

Outcomes	Anticipated absolute effects* (95% CI)  Risk with Dry needling		№ of participants (Studies)	Quality of the evidence (GRADE)	Comments
Pain reduction assessed with: VAS	The mean pain reduction in the intervention group was 14.09 WMD higher (-5.81 lower to 33.99 higher)	-	(4 RCTs)	⊕⊕○○ LOW 12	No significant reduction in pain

<sup>\*</sup>The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).

CI: Confidence interval; RR: Risk ratio; OR: Odds ratio;

#### **GRADE Working Group grades of evidence**

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

- 1. Stor heterogenitet
- 2. Få studier, få deltakere

**Oppsummering:** Resultatene av fire RCT studier viser ingen signifikant smertereduksjon ved tørrnåling sammenlignet med placebo. Denne dokumentasjonen er vurdert å være lav kvaliteten. Grunnlaget for dokumentasjonen er basert på studier med få deltakere og stor heterogenitet.

#### **CAPSAICIN**

Winocur E, Gavish A, Halachmi M, Eli I, Gazit E. Topical application of capsaicin for the treatment of localized pain in the temporomandibular joint area. Journal of Orofacial Pain 2000;14:31–6.

## Topical capsaicin compared to placebo for TMD

Patient or population: TMD Intervention: Topical capsaicin Comparison: Placebo

Outcomes			Relative		Quality of the	Comments
	Risk with placebo	Risk with Topical capsaicin	effect (95% CI)	The second second	evidence (GRADE)	
Reducing pain assessed with: VAS follow up: median 3 months	The mean reducing pain in the control group was 0	The mean reducing pain in the intervention group was 0.61 lower (2.45 lower to 1.23 higher)	-	30 (1 RCT)	⊕⊕⊖⊖ LOW 1234	There was no statistically significant difference between topical capsaicin and placebo in reducing the pain of TMD.

<sup>\*</sup>The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI). CI: Confidence interval: RR: Risk ratio: OR: Odds ratio:

#### **GRADE Working Group grades of evidence**

High quality: We are very confident that the true effect lies close to that of the estimate of the effect

Moderate quality: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low quality: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

Very low quality: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

- 1. Adequate sequence generation?
- 2. Allocation concealment?
- 3. Incomplete outcome data addressed
- 4. Only one study, few participants

**Oppsummering:** Resultatene av en RCT studie viser ingen signifikant smertereduksjon ved bruk av capsaicin sammenlignet med placebo. Dokumentasjonen er vurdert å være av lav kvalitet. Grunnlaget for dokumentasjonen er basert på en studie med få deltakere og stor heterogenitet.