MEDIKAMENTELL BEHANDLING: PARACETAMOL OG NSAID

Roelofs PDDM, Deyo RA, Koes BW, Scholten RJPM, van Tulder MW. Non-steroidal anti-inflammatory drugs for low back pain. The Cochrane Library 2008, Issue 1

Comparison NSAIDs and placebo for Low back pain

Patient or population: Low back pain

Intervention: NSAIDs Comparison: Placebo

Outcomes	Anticipated absolute effects*(95% CI)	Relative	Nº of	Quality of the	Comments
	Risk with NSAIDs	effect (95% CI)	participants (Studies)	evidence (GRADE)	
Change in pain intensity assessed with: VAS follow up: <=3 weeks	The mean change in pain intensity in the intervention group was 8.39 lower (12.68 lower to 4.1 lower)	-	745 (4 RCTs)	⊕⊕⊕○ MODERATE 12	Statistically significant effects in favor of NSAIDs compared to placebo for population with low back pain.
Side effects follow up: <=3 months	The mean side effects in the intervention group was 1.35 RR higher (1.35 higher to 1.68 higher)	-	1852 (10 RCTs)	⊕⊕⊕○ MODERATE 12	Statistically significant side effect in favor of NSAIDs compared to placebo
Proportion of patients experiencing global improvement. Follow-up <=3 weeks.	The mean new outcome in the intervention group was 1.19 RR higher (1.07 higher to 1.33 higher)	-	954 (7 RCTs)	⊕⊕⊕○ MODERATE 12	Statistically significant side effect in favor of NSAIDs compared to placebo

^{*}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. Randomization procedure?
- 2. Allocation concealment?

Summary of findings:

Comparison NSAID to Paracetamol for Low back pain

Patient or population: Low back pain

Intervention: NSAID
Comparison: Paracetamol

			Relative		Quality of the	Comments
	Risk with Paracetamol	Risk with NSAID	effect (95% CI)	participants (Studies)	evidence (GRADE)	
Pain intensity on various scales follow up: mean <=3 weeks	-	The mean pain intensity on various scales in the intervention group was 21 standard deviations lower (0.43 lower to 0.02 higher)	-	309 (3 RCTs)	⊕⊕○○ LOW 1234	There is moderate evidence that NSAIDs are equally effective for pain relief compared with paracetamol for acute low back pain
Proportion of patients side effects follow up: mean <=3 months	-	The mean proportion of patients side effects in the intervention group was 1.76 RR higher (1.12 higher to 2.76 higher)	-	309 (3 RCTs)	⊕⊕○○ LOW 1234	NSAIDs were associated with more side effects compared to paracetamol
Patients experiencing global improvement follow up: <=3 weeks	-	The mean patients experiencing global improvement in the intervention group was 1.23 standard deviations higher (0.88 higher to 1.73 higher)	-	128 (3 RCTs)	⊕⊕○○ LOW 145	There is moderate evidence that NSAIDs are equally effective for global improvement compared with paracetamol for acute low back pain

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- 1. Allocation concealment?
- 2. Patients/Care providers blinded?
- 3. Outcomes blinded?
- 4. Few participants
- 5. Withdrawals

NSAID selective COX-2 inhibitors compared to NSAID non selective COX-2 inhibitors for Low back pain

Patient or population: Low back pain

Intervention: NSAID selective COX-2 inhibitors
Comparison: NSAID non selective COX-2 inhibitors

Outcomes	Anticipated absolute effects*(95% CI)				Quality of the	Comments
	Risk with NSAID non selective COX-2 inhibitors	Risk with NSAID selective COX-2 inhibitors	effect (95% CI)	participants (Studies)	evidence (GRADE)	
Change in Pain Intensity	-	The mean change in Pain Intensity in the intervention group was 2 higher (1.92 lower to 5.92 higher)	-	440 (3 RCTs)	⊕⊕⊕⊕ ніgн	No statistically significant differences for pain relief for acute low-back pain
Proportion of patients experiencing side effects.		The mean proportion of patients experiencing side effects. in the intervention group was 0.83 RR higher (0.7 higher to 0.99 higher)	-	1059 (RCTs)	⊕⊕⊕○ MODERATE ¹	Selective Cox-2 had statistically significantly fewer side- effects compared to Non selective Cox inhibitors

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

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

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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. Allocation, randomization?

Oppsummering: Resultatene viser:

- 1. Signifikant smertereduksjon ved bruk av NSAID sammenlignet med placebo hos pasienter med kroniske lave ryggsmerter. Moderat dokumentasjonskvalitet.

 Bruk av NSAID er like effektivt som bruk av paracetamol for reduksjon av smerter hos pasienter med akutte lave ryggsmerter. NSAID forårsaket mer bivirkninger enn paracetamol. Lav dokumentasjonskvalitet.
- 2. Det finnes ikke statistisk signifikant forskjell i smertereduksjon mellom COX-2 NSAID og «tradisjonelle» NSAID hos pasienter med akutte lave ryggsmerter. Høy dokumentasjonskvalitet.
- 3. COX-2 NSAID forårsaket mindre bivirkninger enn tradisjonelle NSAID hos pasienter med akutte lave ryggsmerter. Moderat dokumentasjonskvalitet.

Dokumentasjonen er vurdert å være av høy, moderat til lav kvalitet.

OBS! Forskningsgrunnlaget er basert på studier av lave ryggsmerter og kan ikke overføres til TMD uten forbehold.