

Summary of findings:




22. Topical analgesia as an adjunct to simple oral analgesia compared to placebo ear drops for immediate pain relief in acute otitis media

Patient or population: Children aged 3 to 19 years with acute otitis media and pain.

Setting: Emergency departments

Intervention: Topical analgesic ear drops as an adjunct to simple oral analgesia (Studies used: 2% aqueous lignocaine or antipyrine benzocaine glycerine. Both studies offered paracetamol 15mg/kg/dose single dose). Single dose of ear drops given on presentation to emergency department.

Comparison: Placebo ear drops (Studies used: Normal saline or olive oil. Both studies offered paracetamol 15mg/kg/dose single dose)

Outcome № of participants (studies)	Relative effect (95% CI)	Anticipated absolute effects (95% CI)			Quality	What happens
		Without Topical analgesia as an adjunct to simple oral analgesia	With Topical analgesia as an adjunct to simple oral analgesia	Difference		
50% reduction in ear pain assessed: two visual analogue scales follow up: median 10 minutes № of participants: 117 (2 RCTs) ^{1,a}	RR 2.13 (1.19 to 3.80)	20.3%	43.3% (24.2 to 77.3)	23.0% more (3.9 more to 56.9 more)	 LOW ^{b,c}	Children with AOMwoP who have local anaesthetic ear drops administered by a health professional compared to placebo possibly have a reduction in pain score by 50% at 10 minutes. NNT ~5.
50% reduction in ear pain assessed: two visual analogue scales follow up: median 20 minutes № of participants: 117 (2 RCTs) ^{1,a}	RR 1.24 (0.88 to 1.74)	47.5%	58.8% (41.8 to 82.6)	11.4% more (5.7 fewer to 35.1 more)	 LOW ^{b,c}	Children with AOMwoP who have local anaesthetic ear drops administered by a health professional compared to placebo possibly have no reduction in pain score by 50% at 20 minutes. NNT Not Applicable
50% reduction in ear pain assessed: two visual analogue scales follow up: median 30 minutes № of participants: 117 (2 RCTs) ^{1,a}	RR 1.43 (1.12 to 1.81)	59.3%	84.8% (66.4 to 100.0)	25.5% more (7.1 more to 48.1 more)	 LOW ^{b,c}	Children with AOMwoP who have local anaesthetic ear drops administered by a health professional compared to placebo possibly have a reduction in pain score by 50% at 30 minutes. NNT ~4

*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; NS: Not significant; NNT: Number needed to treat; NNH: Number needed to harm; AOMwoP: Acute otitis media without perforation

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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

Explanations

- a. Studies taken from: Cochrane Review, Foxlee 2008 (Bolt 2008, Hoberman 1997).
- b. Indirectness: Studies of children 3-18 years of age. Peak incidence AOM is 6-15 months of age.
- c. Imprecision: Small studies

References

1. Foxlee R, Johansson A, Wejfk J, Dawkins J, Dooley L, Del Mar C. Topical analgesia for acute otitis media. The Cochrane database of systematic reviews. 2006(3):Cd005657. Epub 2006/07/21. doi: 10.1002/14651858.CD005657.pub2. PubMed PMID: 16856108.