

Effectiveness of Nebulized Dexmedetomidine in Managing Post-Dural Puncture Headache in Obstetric Patients: A Systematic Review

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Background & Hypothesis

The Clinical Problem

Post-dural puncture headache (PDPH) is a debilitating complication driven by CSF leakage (1).

The Gold Standard: Epidural Blood Patch (EBP)

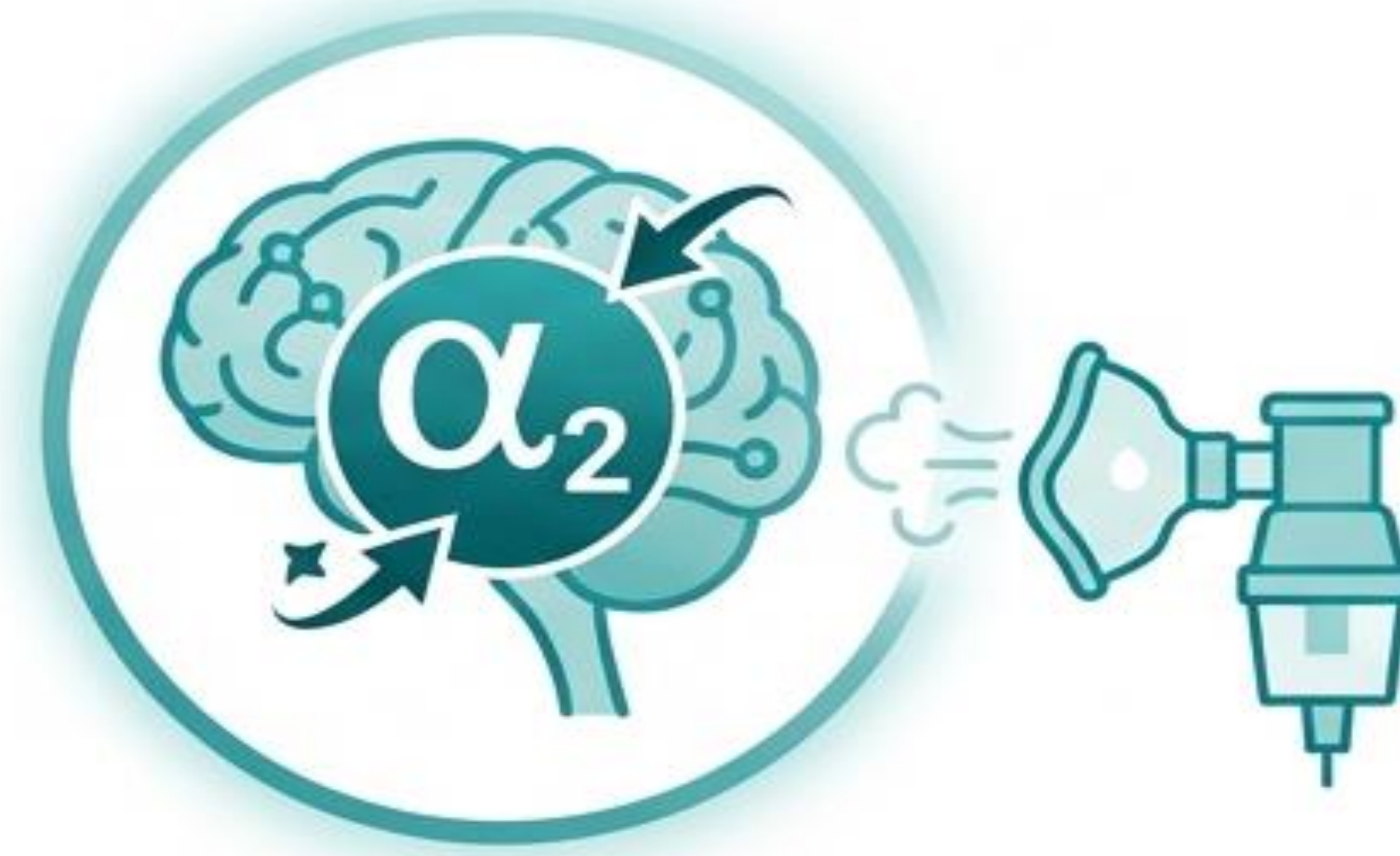
Risks: Back pain, radicular symptoms, bradycardia, meningeal irritation (1,2).



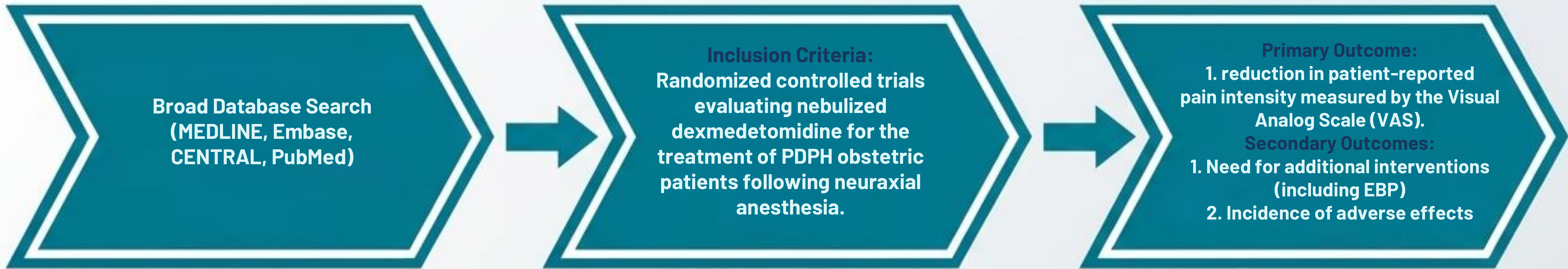
Hypothesis

Nebulized dexmedetomidine reduces PDPH severity and may serve as a non-invasive alternative.

Mechanism: An α_2 -adrenergic agonist providing systemic analgesia, sedation, and cerebral vasoconstriction. (3)

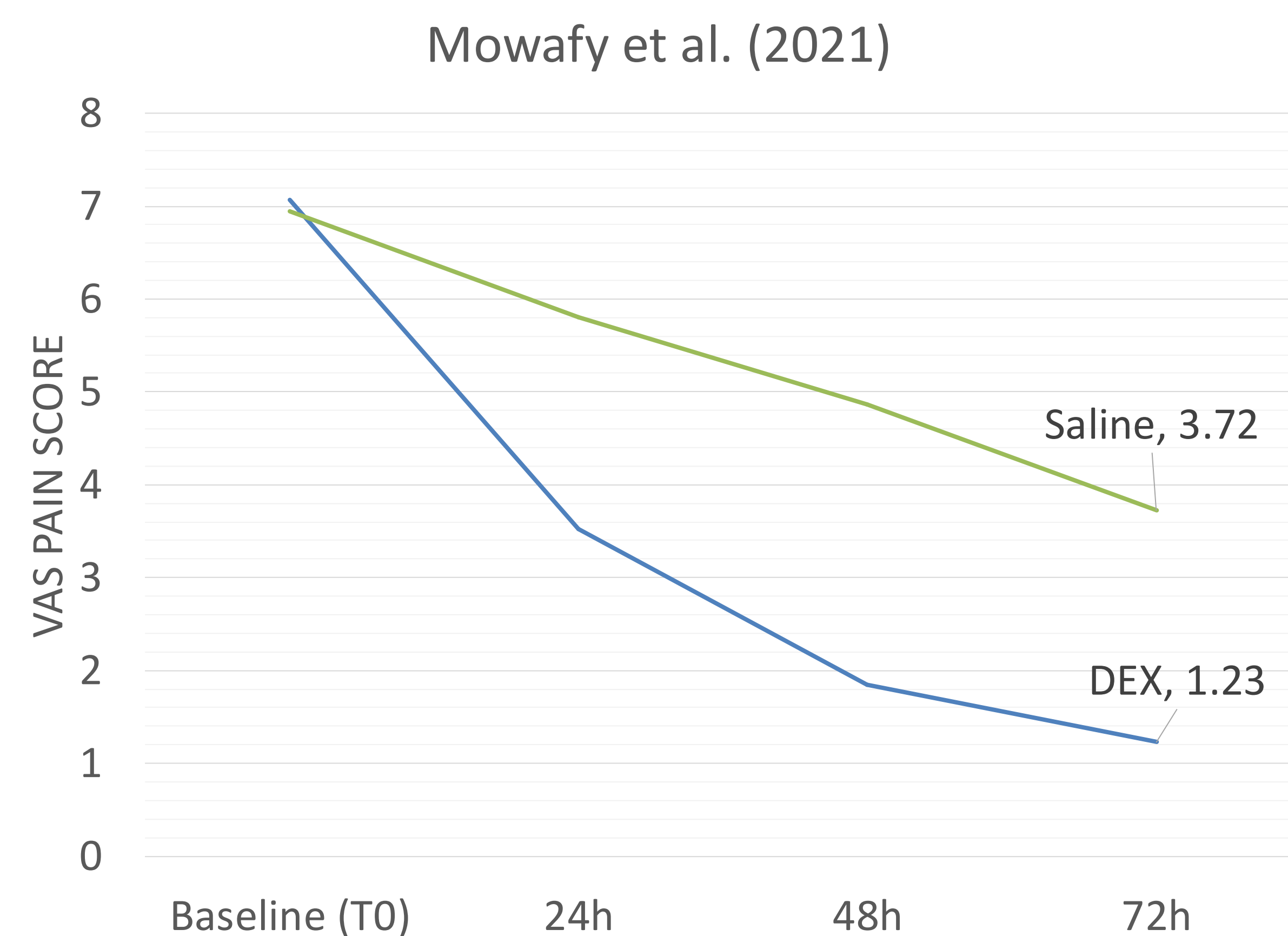
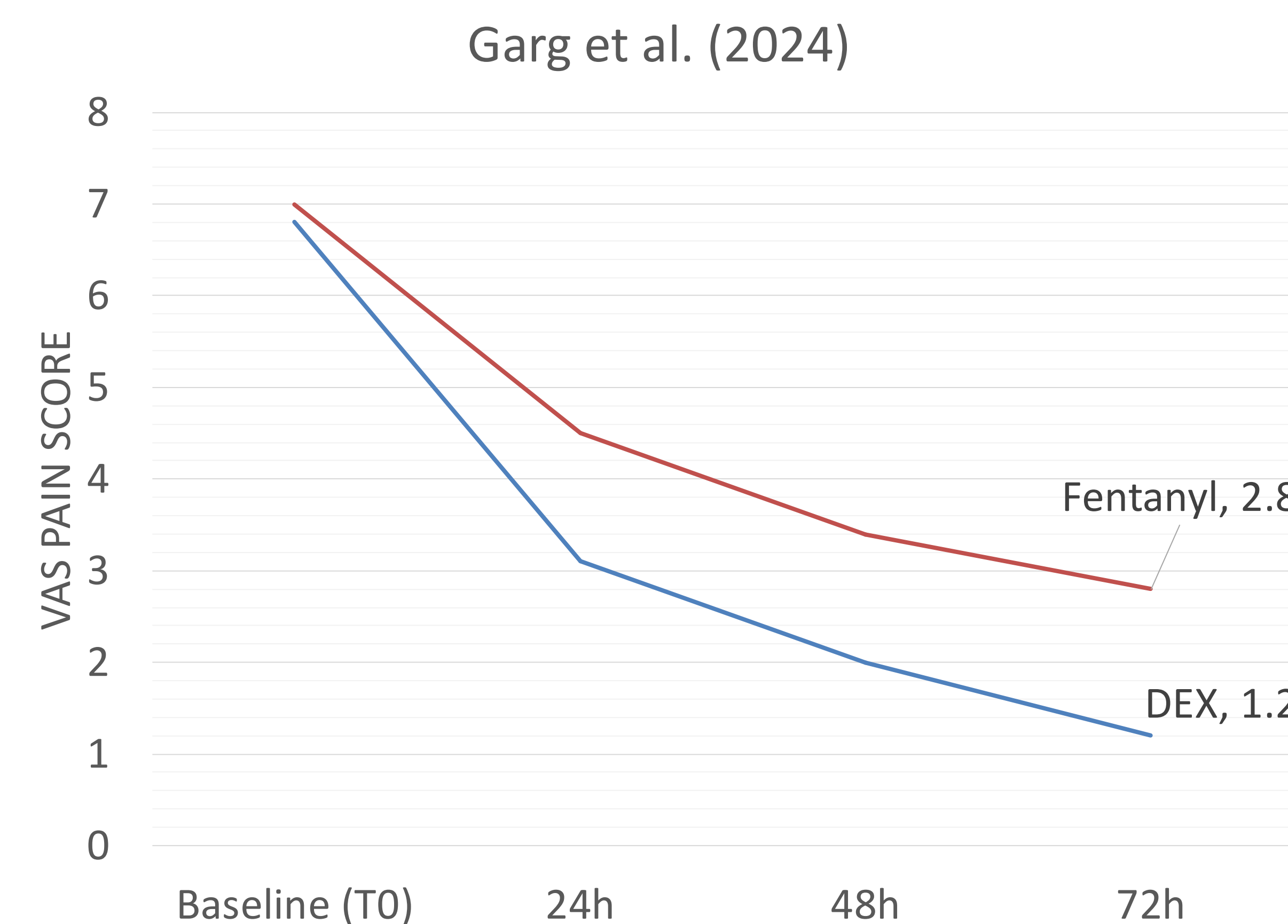
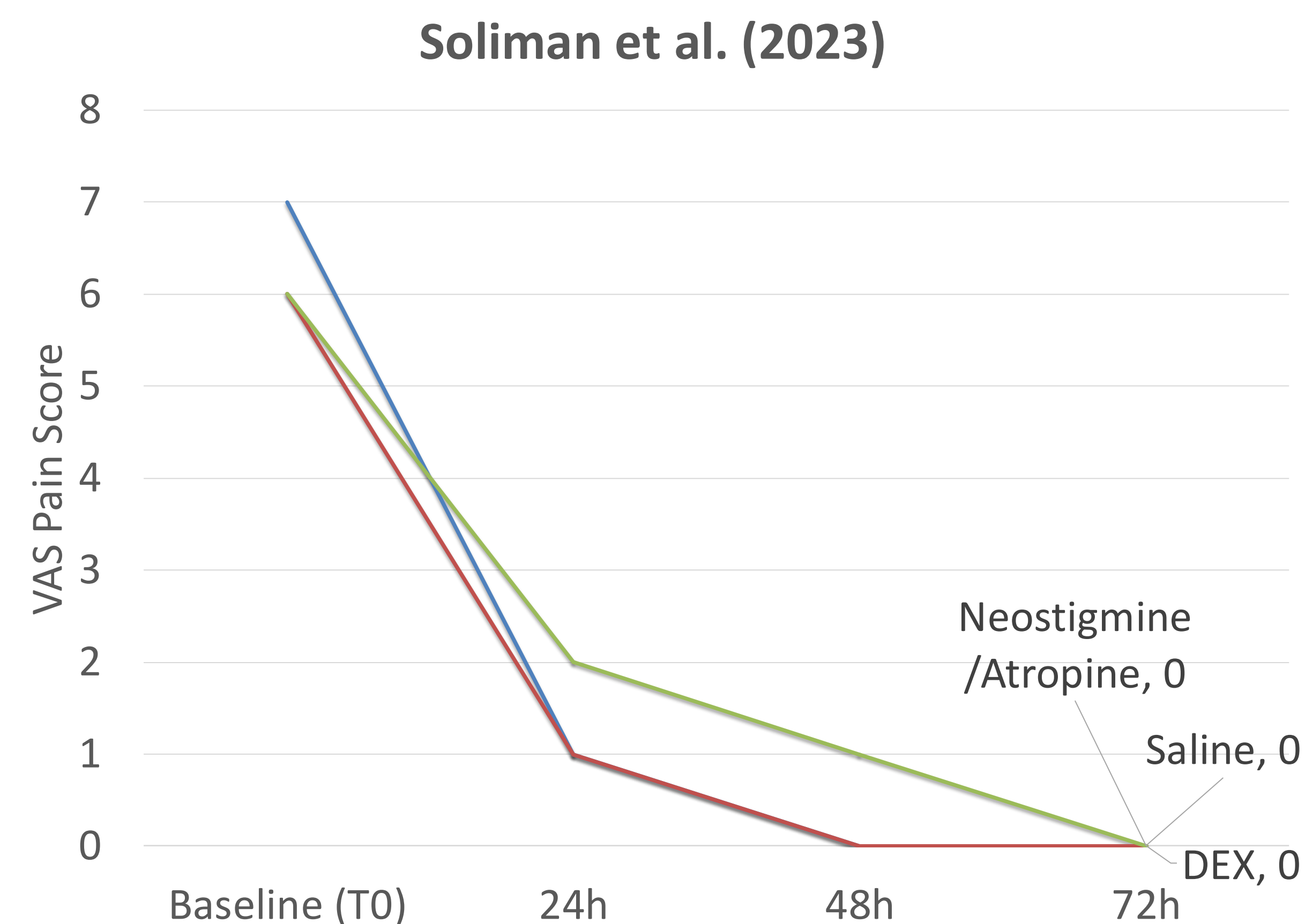
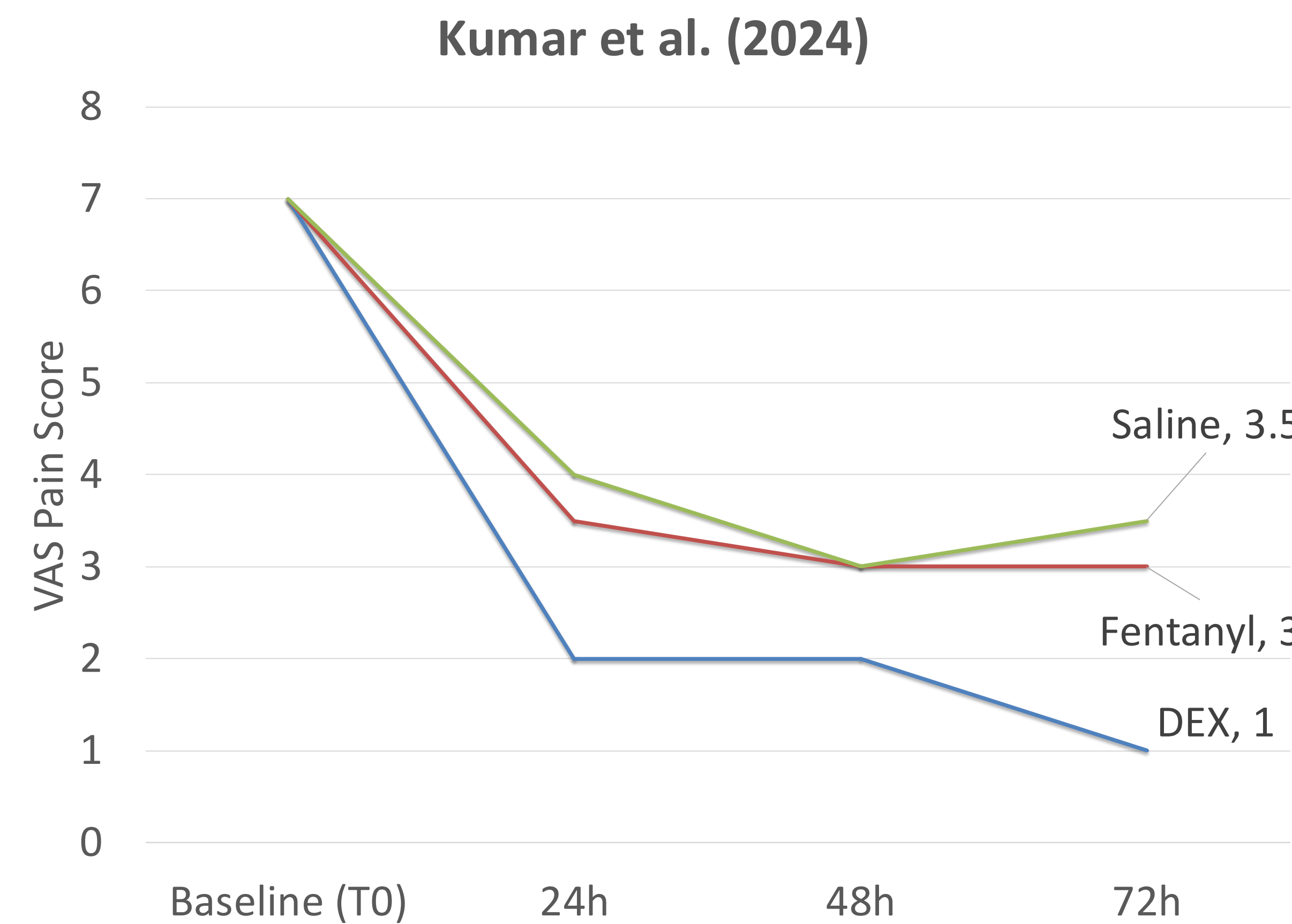


Study Designs & Methods



Study & Location	Patients	Intervention Dose	Comparators	Outcomes Tracked
Kumar et al. (4), India	n=90	1 µg/kg BID x 3 days	Fentanyl, Saline	VAS (24/48/72h), Adverse Events, EBP needed
Soliman et al. (5), Egypt	n=90	1 µg/kg BID x 3 days	Neostigmine/Atropine, Saline	VAS (24/48/72h), Adverse Events, EBP needed
Garg et al. (6), India	n=65	Not specified.	Fentanyl	VAS (24/48/72h), Adverse Events
Mowafy et al. (3), Egypt	n=43	1 µg/kg BID x 3 days OR until VAS ≤3 and Lybecker score <2	Saline	VAS (24/48/72h), Adverse Events, EBP needed

Results



Across all RCTs, DEX consistently delivered superior pain control versus placebo and Fentanyl, and comparable with Neostigmine / Atropine.

Rescue Therapy Need

0 EBP

Across all 4 RCTs (n=92), zero patients in the DEX groups required an Epidural Blood Patch.



DEX: Primarily drowsiness/sedation, dry mouth, and bradycardia



Fentanyl: Primarily drowsiness, nausea, dizziness



Neostigmine/Atropine: Diarrhea, abdominal cramps, and muscle twitches

Conclusion & Discussion

Nebulized dexmedetomidine appears to be a safe and effective non-invasive option for obstetric PDPH.

- Provides faster symptom relief to traditional pharmacological options (i.e., fentanyl), and comparable efficacy to neostigmine/atropine.
- Limited need for adjunctive treatments, including no need for EBP.
- Tolerable side effects similar in incidence with comparators.

Clinical Strengths



- Non-invasive administration.
- Consistent efficacy established at 1 µg/kg dose diluted to 4 mL.
- Enhances systemic bioavailability by bypassing first-pass metabolism, no need for intravenous access.

Evidentiary Limitations



- VAS is inherently subjective.
- Off-label status for nebulized administration in North America.
- Evidence is limited to PDPH following spinal anesthesia for LSCS and has not been studied in epidural-related PDPH.

Next Steps: Larger, multicenter trials are needed to validate these findings and to better define its role in the routine prevention and treatment of PDPH.

THANK YOU!



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