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# Injectable In Situ Thermoreversible Gel Depot System of Lidocaine Nanoemulsion for Prolonged Anesthetic Activity in Dental and Operative Procedures

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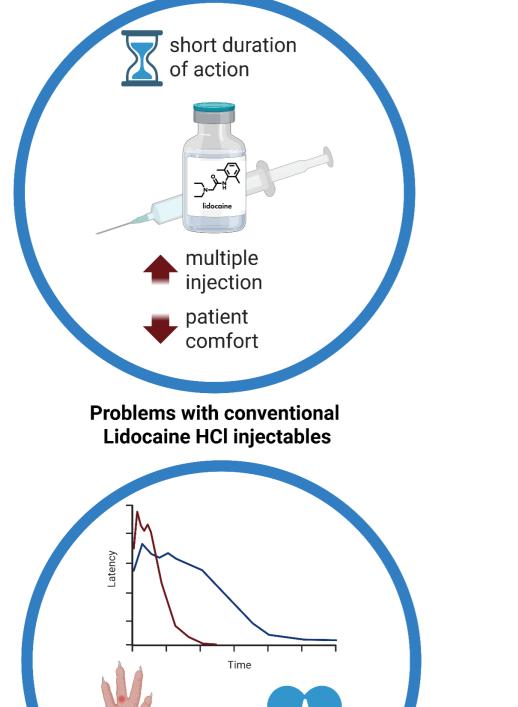
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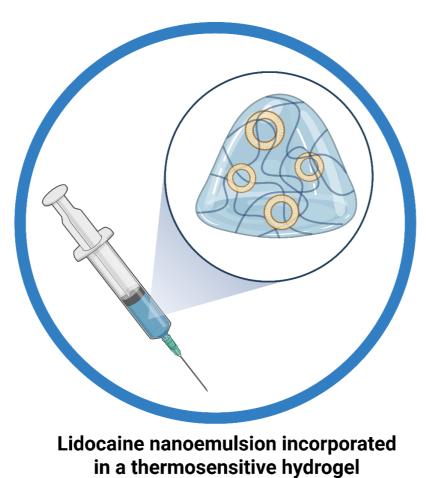
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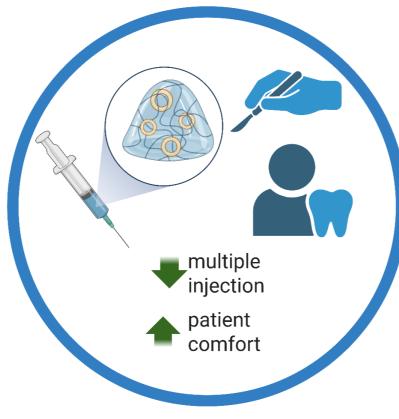
#### INTRODUCTION & AIM

- Lidocaine hydrochloride (LD-HCl) is the most commonly used local anesthetic in dentistry, often administered with epinephrine to extend its duration and reduce systemic absorption.
- However, its relatively short duration of action, the need for repeated injections, and the unpleasant taste may limit patient compliance and procedural efficiency.
- This study aimed to develop and evaluate a novel injectable nanoemulsion-based in situ gel depot system of LD to provide prolonged anesthetic activity.



Prolonged anesthetic activity with no cardiovascular adverse effects

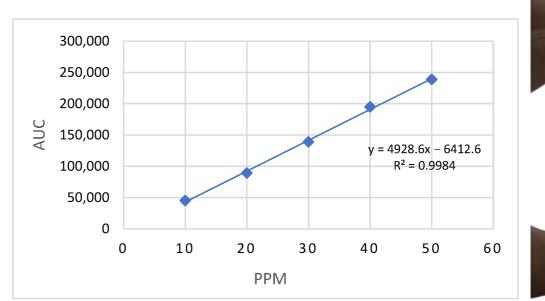


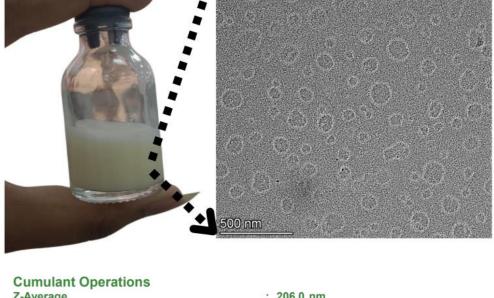


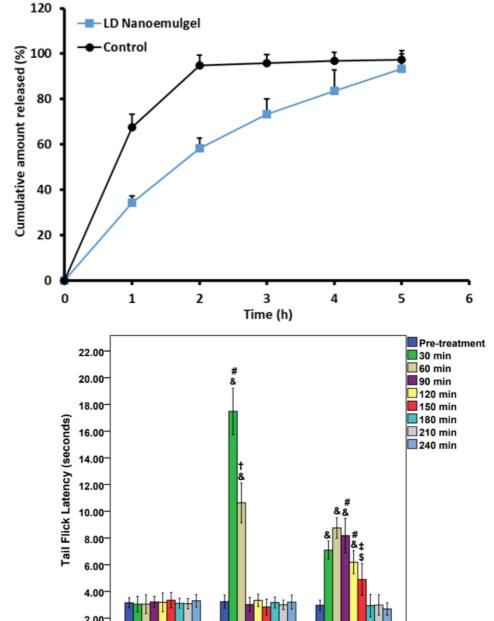
Better anesthetic outcomes for dental and surgical patients

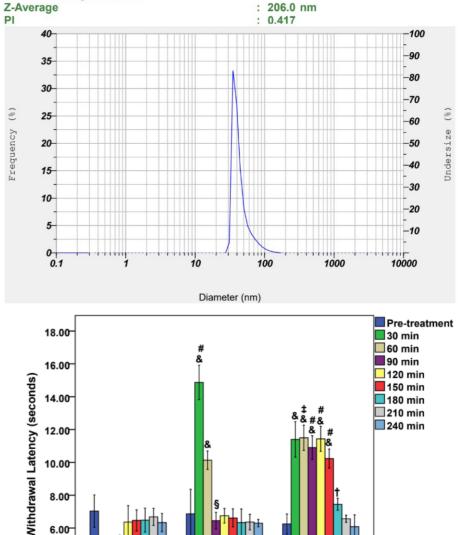
## RESULTS & DISCUSSION

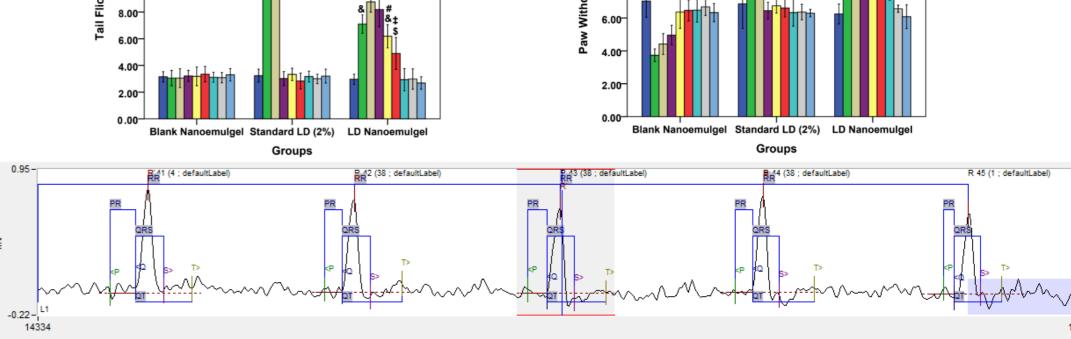
- The developed nanoemulsions demonstrated a spherical shape, nanometer size (206 nm), high zeta-potential (-66.67 mV) and uniform size distribution, with a polydispersity index of approximately 0.40, while the nanoemulgel demonstrated appropriate thixotropic properties for parenteral administration.
- In vitro release profiles showed steady LD release (5 h), following the Higuchi model.











# METHOD

- LD-loaded nanoemulsions were formulated by high-shear homogenization followed by probe sonication, employing Miglyol 812 N (oil phase), a combination of Tween 80 and soy lecithin (surfactant-co-surfactant), glycerin (stabilizer), and deionized water (aqueous phase). The selected nanoemulsion was dispersed in a thermoreversible poloxamer solution to form a nanoemulgel.
- The preparation was evaluated for globule diameter and uniformity, zeta potential, surface morphology, pH, drug content, stability, rheological behavior, injectability, and in vitro drug release.
- Analgesic efficacy was assessed via tail-flick and thermal paw withdrawal latency tests in Wistar rats.
- Cardiovascular safety was monitored using non-invasive electrocardiography and blood pressure measurements.

In vivo studies showed significantly **prolonged analgesic effects lasting up to 150 min (2.5 h)** compared to standard LD-HCl injection (p < 0.001), with **no adverse cardiovascular effects** observed.

#### CONCLUSION

The developed injectable LD in situ nanoemulgel offers a promising, patient-friendly alternative for prolonged anesthetic delivery in dental and operative procedures, potentially reducing the need for repeated injections and enhancing procedural comfort.

### REFERENCES

Link to Publication and references:

Jacob, S., Kather, F. S., Satyam, S. M., Boddu, S. H. S., Assaf, F., Allam, T. H. A., & Nair, A. B. (2025). Injectable In Situ Thermoreversible Gel Depot System of Lidocaine Nanoemulsion for Prolonged Anesthetic Activity in Dental and Operative Procedures. Pharmaceutics, 17(10), 1355. https://doi.org/10.3390/pharmaceutics17101355