Enhanced Viability of HK-2 Cells Encapsulated in Chitosan Hydrogel Compared to Sodium Alginate Hydrogel

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INTRODUCTION

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- Cell encapsulation within hydrogels holds promise for diverse biomedical applications such as tissue engineering and drug delivery. The selection of hydrogel material significantly impacts cell viability and function.
- This study compares the viability of human kidney epithelial cells (HK-2) encapsulated in two hydrogel types: sodium alginate (NaAlg) and chitosan (CS).

METHODS





RESULTS

• The study revealed significantly higher viability of HK-2 cells when encapsulated in CS hydrogels compared to NaAlg hydrogels.





Chitosan hydroge

Sodium alginate hydrogel

- **Fig. 2.** Growth of HK-2 cells in sodium alginate hydrogel and chitosan hydrogel.
- Live/Dead staining indicated a higher percentage of live cells in the CS group, reflecting enhanced metabolic activity.



Fig. 3. AO/PI immunofluorescence staining of HK-2 cells in sodium alginate hydrogel and chitosan hydrogel

RESULTS

 HK-2 cells grew spherically in the hydrogel, and immunofluorescence staining showed significant expression of renal tubular epithelial important marker (Lotus Tetragonolobus Lectin, LTL) in HK-2 rechitosan hydrogel.



Fig. 4 Immunofluorescence staining showed that renal tubules were mainly composed of proximal tubular epithelial cells, and HK-2 cells showed good activity in chitosan hydrogels.

CONCLUSION

- These findings highlight the potential of CS hydrogels as excellent scaffolds for HK-2 cell inclusion, with important implications for renal tissue engineering strategies and applications needed to maintain renal cell function.
- Chitosan hydrogels were better than sodium alginate hydrogels in 3D culture of HK-2 cells to maintain cell activity and proximal tubular epithelial characterization
- Future studies should delve into the mechanisms that enhance activity and optimize CS hydrogel components for specific biomedical uses.