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New Formulation of Platelet-Rich Plasma Enriched in Platelet and Extraplatelet Biomolecules Using Hydrogels

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Faculty Disclosure Information

Nothing to disclosure



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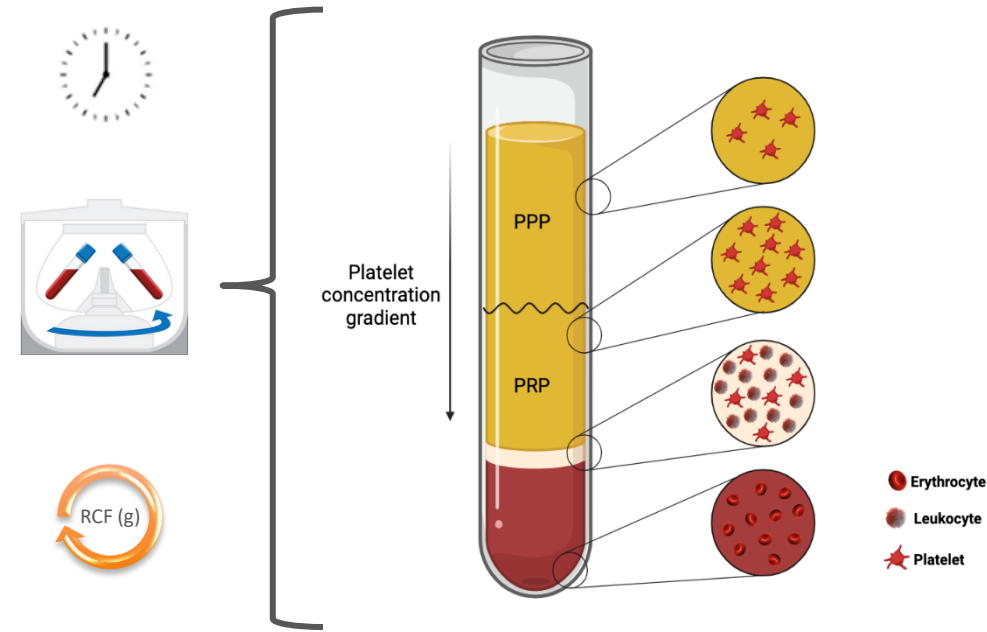


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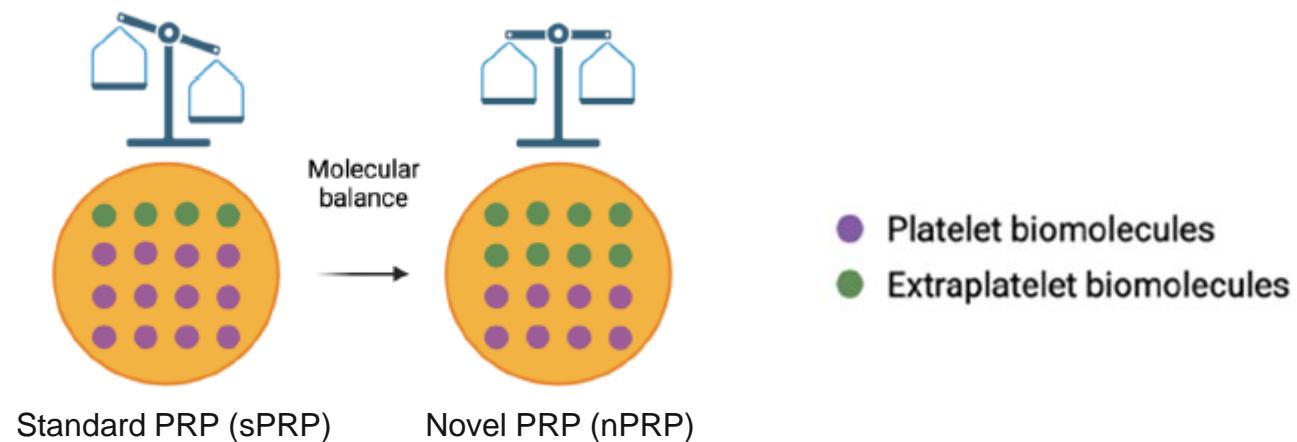
Introduction

- **Platelet-rich plasma (PRP)** is an **autologous biologic product used in several fields of medicine** for tissue repair due to the regenerative capacity of the biomolecules of its formulation.
- PRP consists of a plasma with a **platelet concentration higher than basal levels** but with basal levels of any biomolecules present out of the platelets
- Conventional methods of obtaining PRP are unable to modulate **extraplatelet molecules**.



Objective

- The aim of this work is to develop a new method to obtain **PRP enriched in both platelet and extraplatelet molecules**.
- The hypothesis of this study is based on the fact that PRP containing not only a higher concentration of platelets but also **a higher concentration of extraplatelet biomolecules** could have a **stronger regenerative performance than a standard PRP**.



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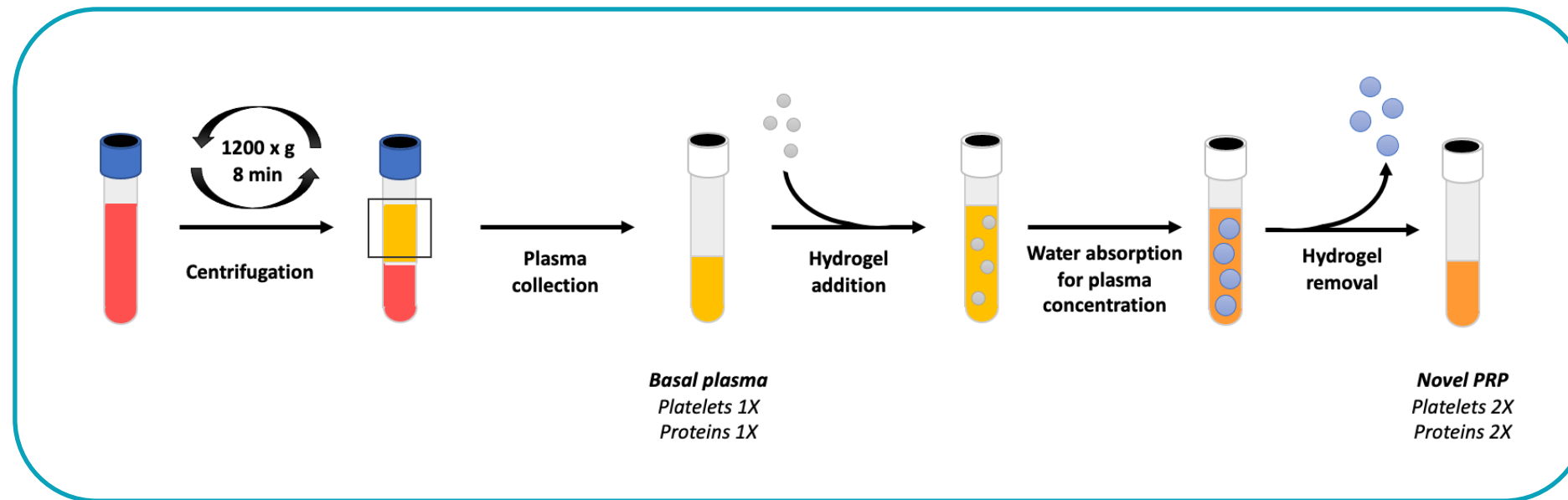


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Methods

- A **plasma fraction** obtained from blood, containing the basal levels of platelets and proteins, was placed in contact with the **HEAA hydrogel powder** to **absorb half the volume of the water**



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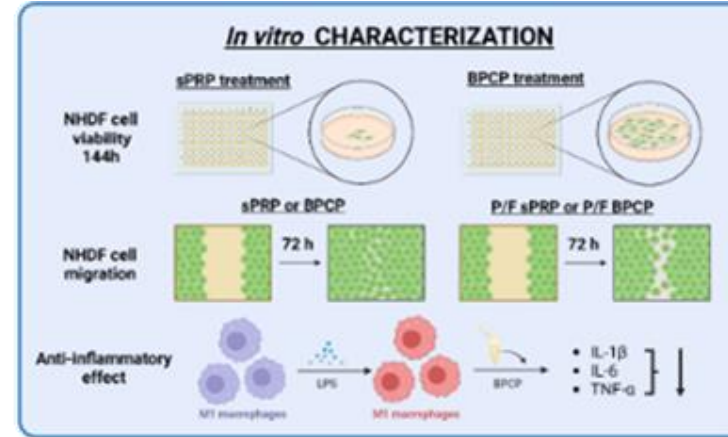
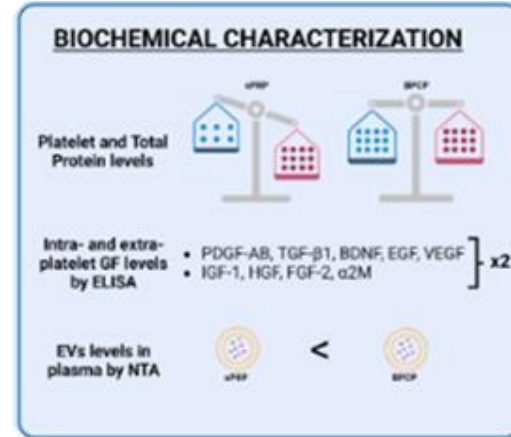
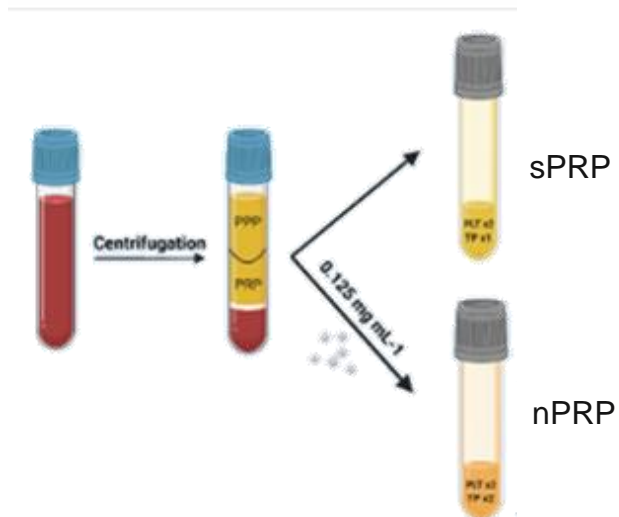


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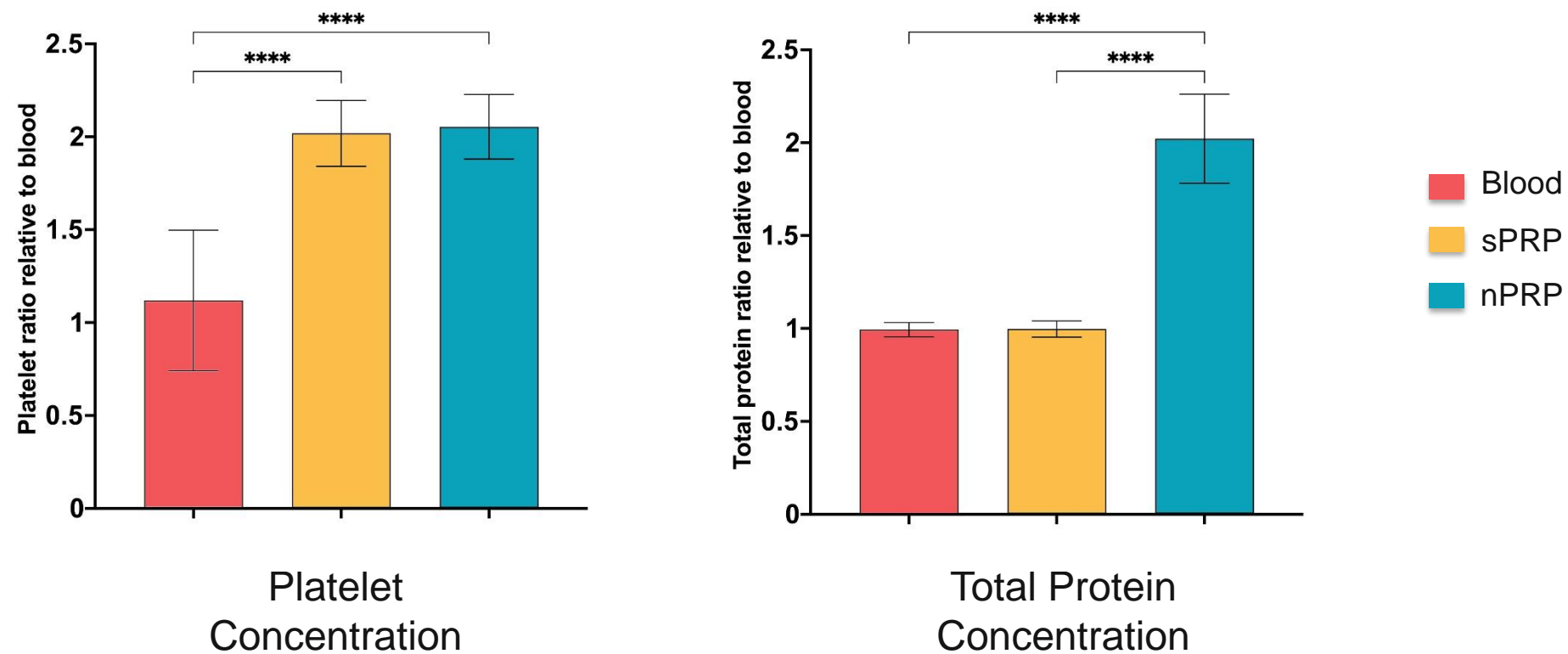
Methods

- The resulting plasma (nPRP) was characterized, and its bioactivity was analyzed in vitro.
- Standard PRP (sPRP) was used as a **control**.



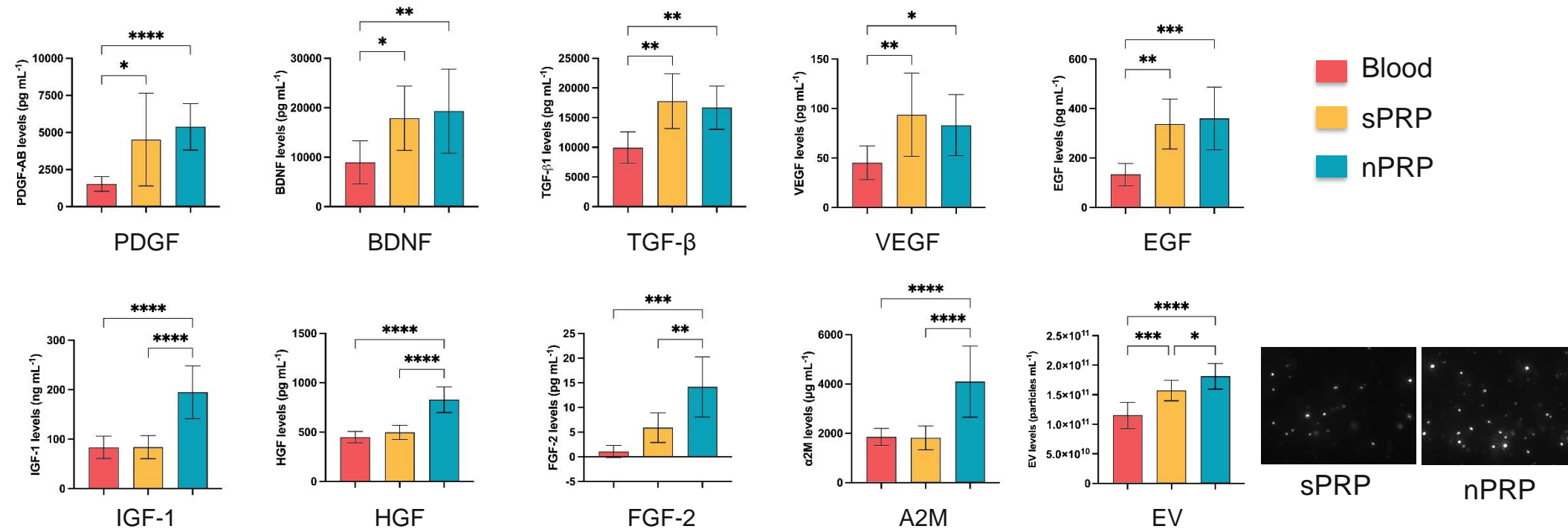
Results - Characterization

The novel PRP (nPRP) showed a **platelet concentration similar** to the standard PRP (sPRP) ($p>0.05$), but the concentration of **the total proteins were significantly increased** ($p<0.0001$).



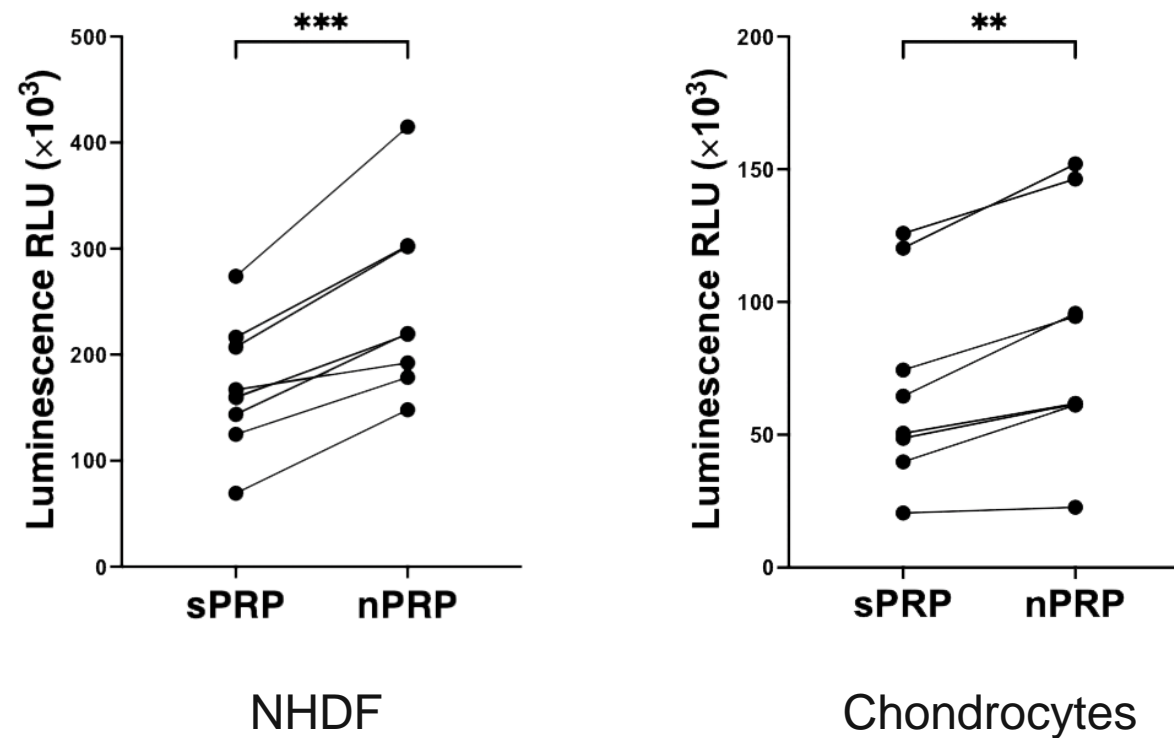
Results - Characterization

- Platelet factors had the same levels in both PRP ($p>0.05$): PDGF, BDNF, TGF- β , VEGF, EGF.
- nPRP had higher levels of extraplatelet molecules and extracellular vesicles (EV) ($p<0.05$): IGF-1, HGF, FGF-2, A2M.



Results – Cell viability

The cells exposed to the **nPRP showed increased cell viability** than those exposed to a sPRP in human dermal fibroblasts (NHDF) ($p < 0.001$) and primary chondrocytes ($p < 0.01$)



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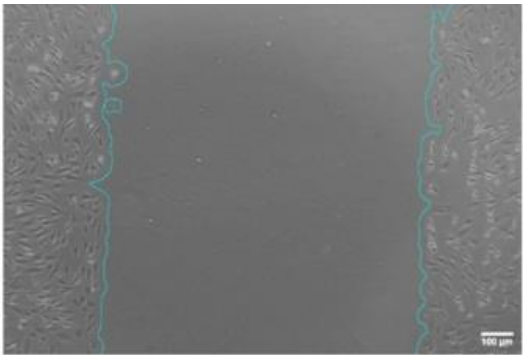
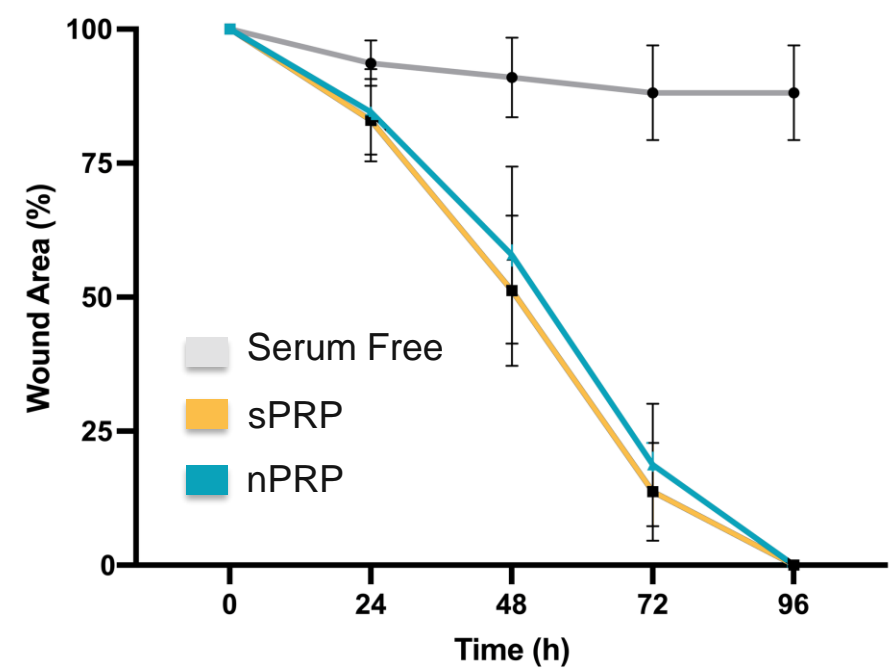
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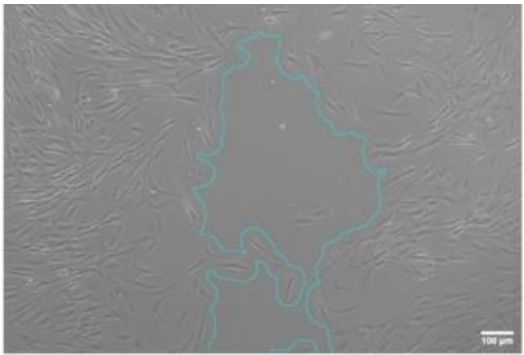
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Advanced Biological Therapy Unit

Results – Cell migration

Regarding to **cell migration capacity**, it was found that the process is platelet-dependent, achieving **similar wound healing closure percentage between sPRP and nPRP**.



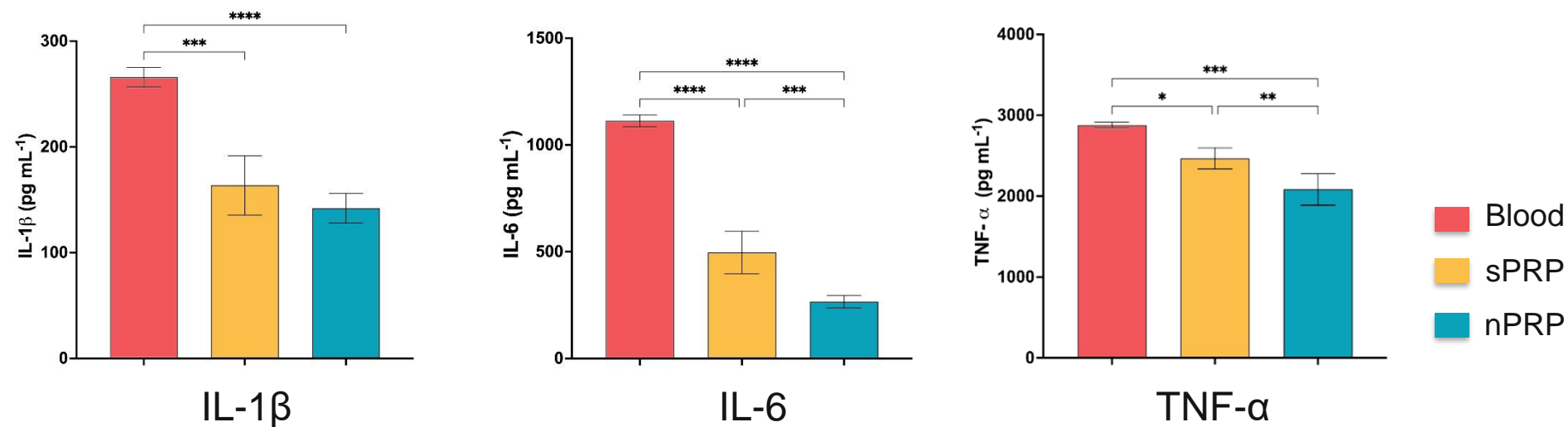
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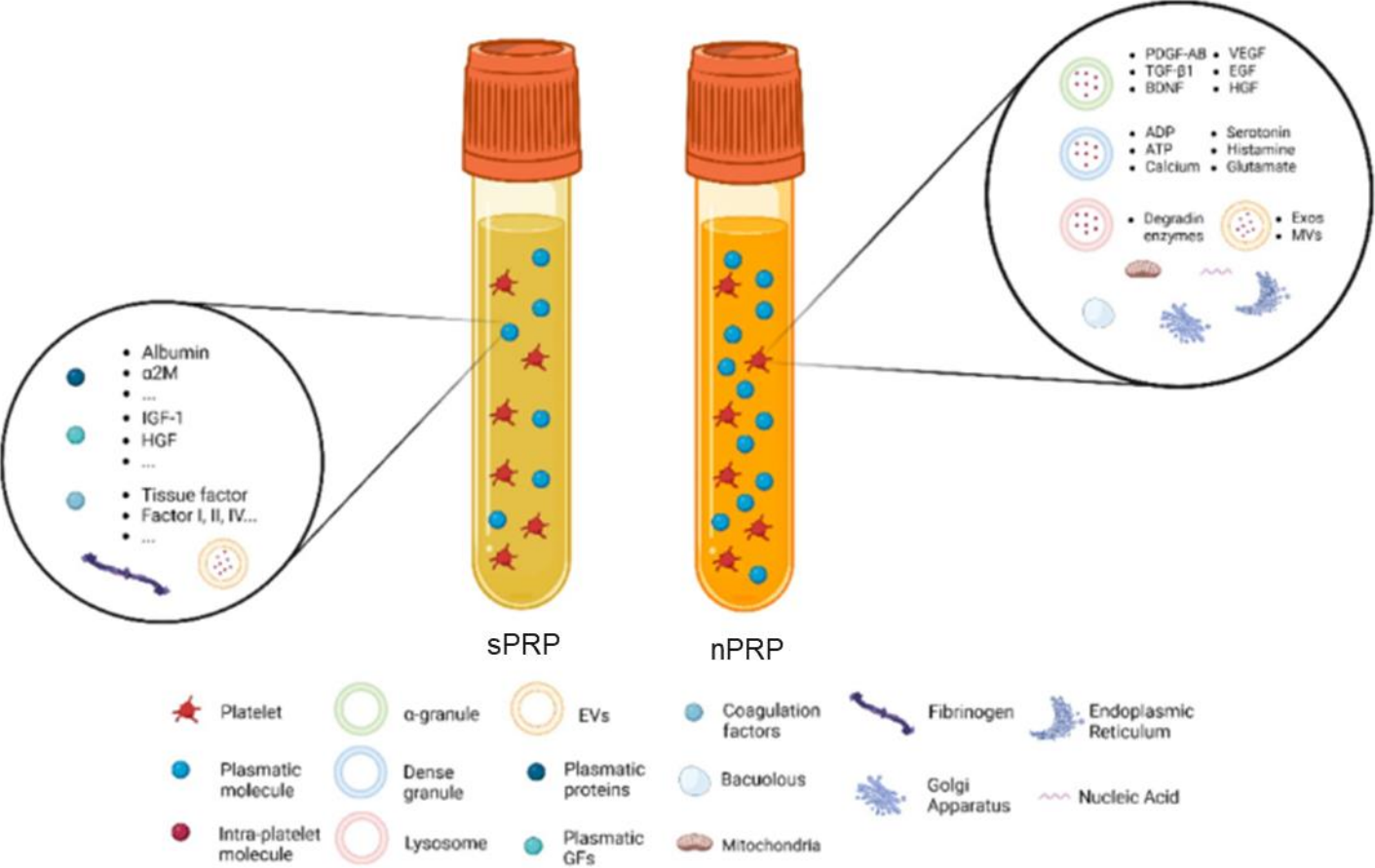
Results - Inflammation

- An **inflammatory environment** was produced using **LPS** in previously differentiated **M1 macrophages**.
- When cytokine levels were measured by ELISA after **nPRP** administration, a significant **decrease of pro-inflammatory IL-1b** ($p < 0.0001$), **IL-6** ($p < 0.0001$) and **TNF- α** ($p < 0.001$) was observed.



Conclusion

This novel absorption-based method produces a PRP with novel characteristics compared to the standard PRPs, with promising in vitro results that could potentially trigger improved tissue regeneration capacity.



References

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Increasing the concentration of plasma molecules improves the biological activity of platelet-rich plasma for tissue regeneration

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