XXVI Congresso de Iniciação Científica Unicamp

17 a 19 de outubro

Campinas | Brasil

Nitric oxide releasing collagen membranes for the topical treatment of wounds

Giovanna J. V. P. dos Santos*; Guilherme F. Picheth; Valéria Póvoa; Eliana P. de Araujo; Marcelo G. de Oliveira.

Abstract

Nitric oxide (NO) donor biomaterials have great potential for the promotion of tissue regeneration. Collagen membranes impregnated with S-nitrosoglutathione (GSNO), an NO donor, were developed, characterized and applied in vivo, demonstrating the acceleration of wound healing.

Key words:

biomaterials, collagen, nitric oxide

Introduction

The use of dressings derived from collagen is already established in the medical area for the coating of lesions and healing aid. Nitric oxide (NO) is directly involved in angiogenic and proliferative activities, stimulating the tissue repair process in healthy individuals or patients suffering from chronic diseases (e.g. diabetes). The main objective of this project is the development of type I collagen membranes functionalized with S-nitrosoglutathione (GSNO) for the localized release of NO.

Results and Discussion

Collagen membranes were characterized by FTIR, SEM/EDS. The NO release of the membranes was characterized by chemiluminescence.

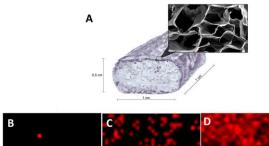


Image 1. (A) Macroscopic photograph of the membrane with details of the internal structure of macropores obtained by SEM. (B, C and D) Sulfur mapping of native collagen (B), collagen with dose 1 of GSNO (C) and dose 2 of GSNO (D) (dose 2>dose 1).

The NO donor collagen membranes displayed high porosity (IMG. 1) and NO release kinetics dependent on the hydration degree (IMG. 2).

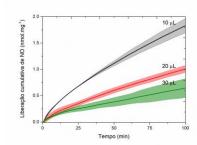


Image 2. Kinetic curves of NO released from collagen/GSNO membranes swollen with 10 μ L, 20 μ L and 30 μ L of PBS with EDTA.

In vivo experiments showed a higher wound healing rate after 4, 6 and 8 days (IMG. 3), as well as larger amounts of cytokines with proliferative activity (e.g. IGF-1 and SDF-1), and overexpression of iNOS and MMP-9 in animals treated with collagen/GSNO membranes.

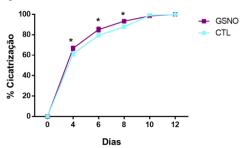


Image 3. Evolution of the lesion area of animals treated with collagen/GSNO membranes (GSNO) and native collagen membrane (CTL), over 12 days.

Conclusions

The collagen membranes impregnated with GSNO accelerated cell proliferation and wound healing possessing potential for the treatment of chronic wounds.

Acknowledgement

GJVPS received a studentship from the São Paulo Research Foundation (FAPESP) (Grant nº 2017/046150). This work was supported by FAPESP (grant number 2016/02414-5).

¹Champeau, M., Póvoa, V., Militão, L., Cabrini, F. M., Picheth, G. F., Meneau, F., Jara, C. P., de Araujo E. P., de Oliveira, M. G. (2018). Supramolecular poly (acrylic acid)/F127 hydrogel with hydration-controlled nitric oxide release for enhancing wound healing. *Acta biomaterialia*, *74*, 312-325

²Shoulders, M. D., & Raines, R. T. (2009). Collagen structure and stability. *Annual review of biochemistry*, 78, 929-958.