

## Participation of microRNA-34a/RANKL in the osteogenic potential of the Poly(vinylidenetrifluorethylene)/barium titanate membrane



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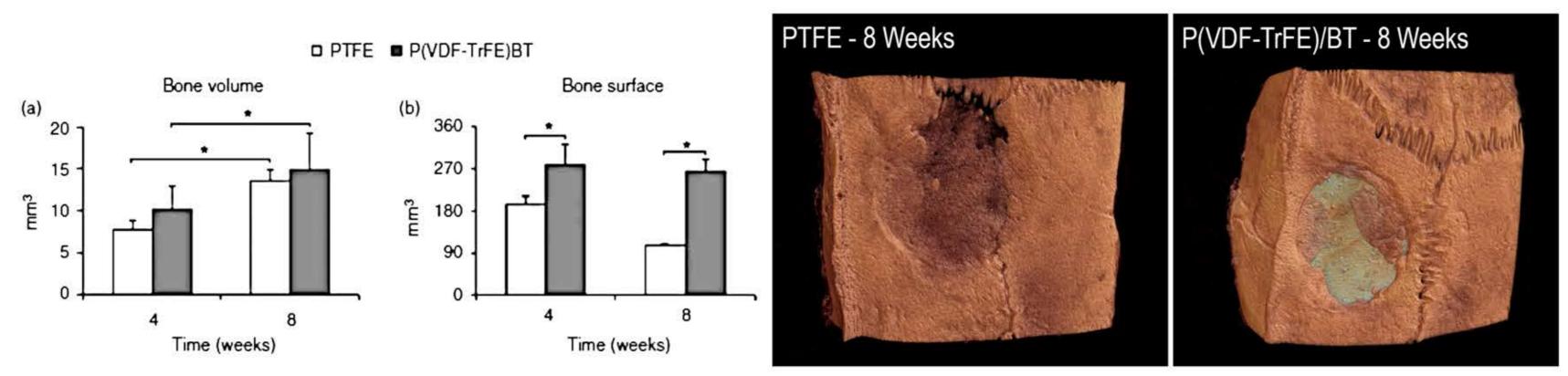
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## **Purpose of Study**

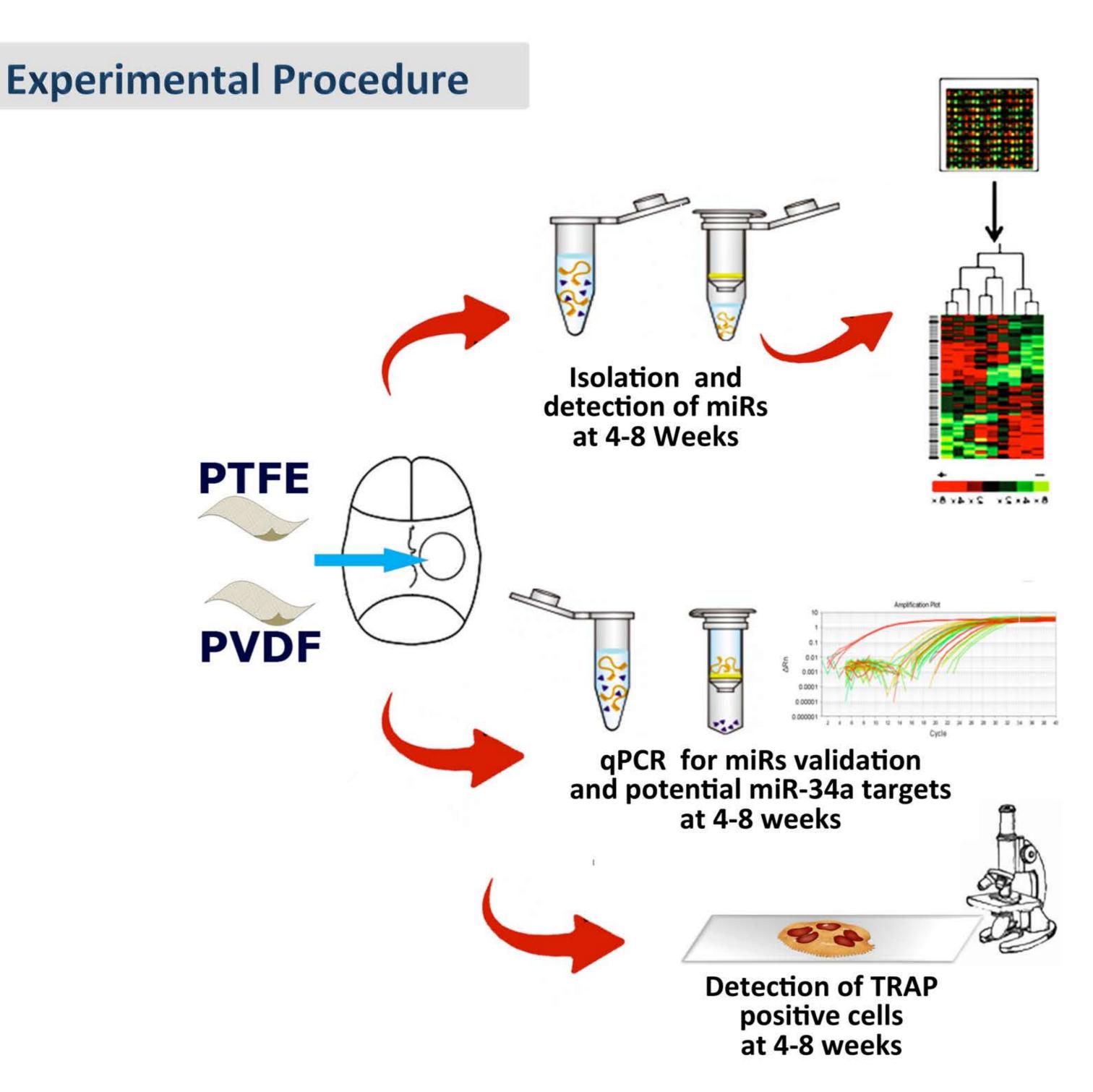
The aim of this study was to investigate a possible mechanism involving miRs and RANKL in the osteogenic potential of the Poly(vinylidene-trifluoroethylene)/barium titanate composite membrane (PVDF).

## Background

Barrier membranes have been extensively used in dentistry to prevent soft tissue down-growth into the bone defects and to promote alveolar ridge augmentation. Previous studies of our group showed that the PVDF enhances both the in vitro osteoblastic differentiation and the in vivo bone repair compared with a commercially available polytetrafluoroethylene (PTFE) membrane.



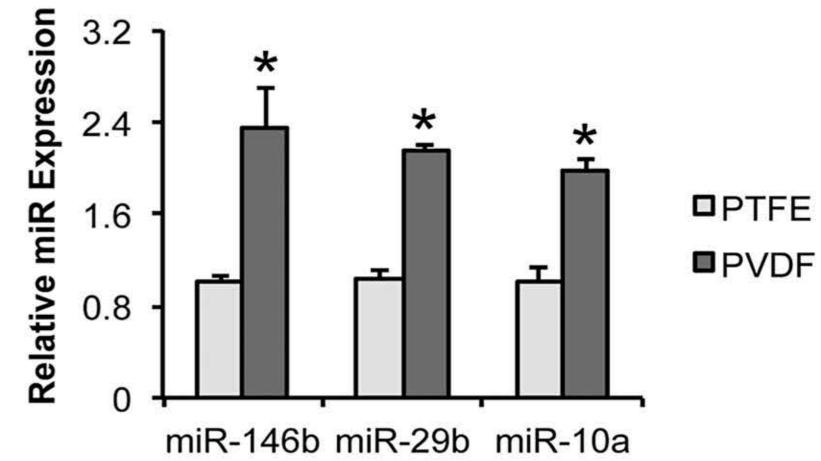
(Lopes et al. Journal of Biomaterials Applications. Vol. 29(1) 104–112. 2014)



# 4 weeks 65 miRs Up/Down 12 miRs Up-regulated miR-146b miR-146b miR-10a miR-10a miR-34a miR-34a miR-29b

**Figure 1.** miR detected at fold change ≥ 2 (up or downregulated) p < 0.05 and q-value < 0.05

### Validation of miR Expression



**Figure 2.** Expression of miR-146b at 4 weeks, and miR-29b and -10a at 8 weeks. \* indicate statistical significant difference ( $p \le 0.05$ ).

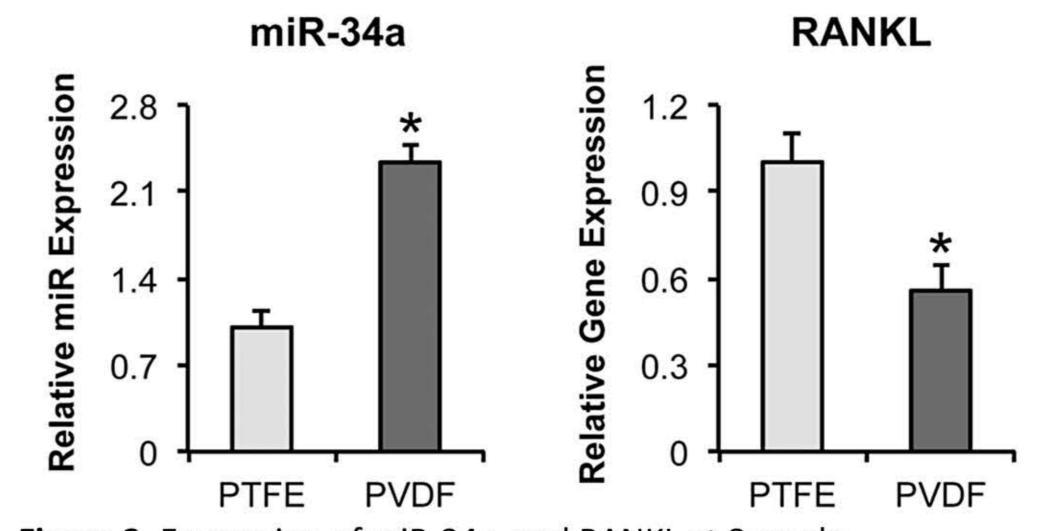


Figure 3. Expression of miR-34a and RANKL at 8 weeks.

\* indicate statistical significant difference (p ≤ 0.05).

# Potential miR-34a Targets 1.6 1.2 0.8 0.4 \* Potential miR-34a Targets PPTFE PPVDF

YY1

SYT1

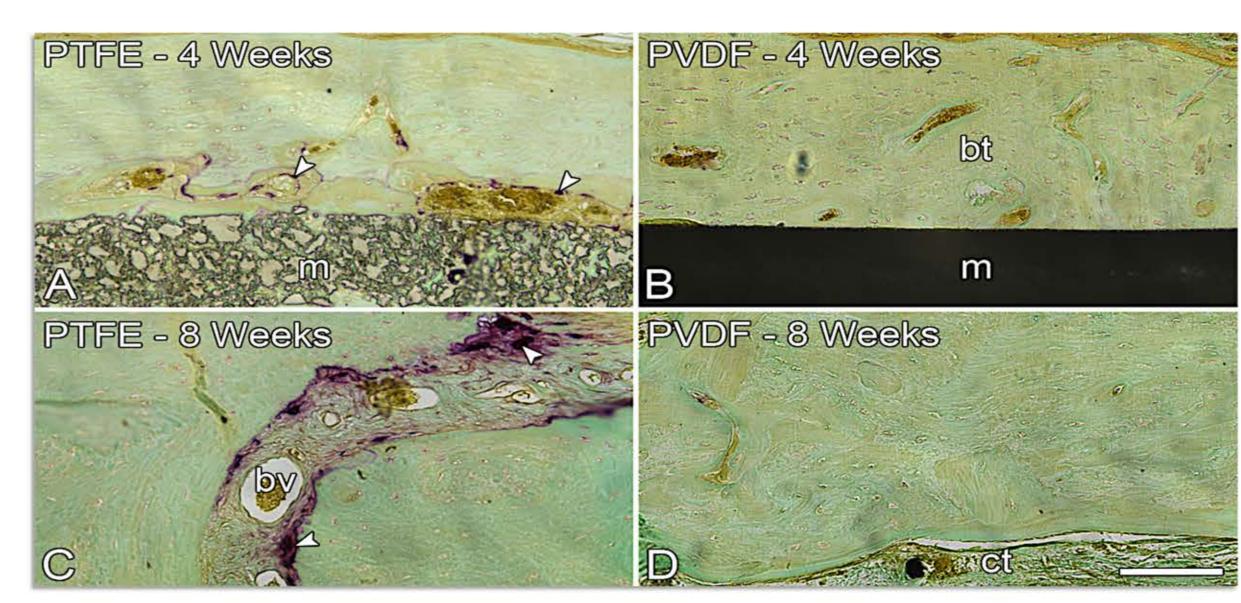
**CSFR** 

Figure 4. Expression of miR-34a targets at 8 weeks. \* indicate statistical significant difference ( $p \le 0.05$ ).

**ENG** 

LEF1

Relative Gene Expression



**Figure 5**. Light microscopy of rat calvarial bone defects implanted with (PTFE; A and C) or (PVDF, B and D) membrane at 4 (A and B) and 8 (C and D). bt: bone tissue; bv: blood vessel; ct: connective tissue; m: membrane. Scale bar:  $A-D = 100 \mu m$ .

### **Concluding Remark**

PVDF membrane induces higher bone repair, at least in part, by triggering an intracellular mechanism of miR-34a upregulation/RANKL downegulation loop, which inhibits osteoclastic activity.