



DOI: <http://dx.doi.org/10.21270/archi.v6i0.2255>

Zer-005

Mineral Trioxide Aggregate improves healing response of periodontal tissue to injury in mice

Queiroz IOA^{1,2}, Vidovic I³, Roeder E², Wang X², Matthews BG², Gomes-Filho JE¹, Mina M³, Kalajzic I²

¹São Paulo State University (UNESP), School of Dentistry, Araçatuba, Department of Restorative Dentistry

²Department of Reconstructive Sciences, Uconn Health Center, Farmington, EUA

³Department of Pediatric Dentistry, Uconn Health Center, Farmington, EUA

Mineral Trioxide Aggregate (MTA) is a biomaterial used in endodontic procedures as it exerts beneficial effects on regenerative processes. In this study, we evaluate the effect of MTA on healing of periodontal ligament (PDL) and surrounding tissue following injury in a transgenic mouse model, and on differentiation of murine mesenchymal progenitor cells *in vitro*. We used an inducible Cre-loxP *in vivo* fate mapping approach to examine the effects of MTA on the contributions of descendants of cells expressing α SMACreERT2 transgene (SMA9+) to the PDL and alveolar bone after experimental injury to the root furcation on the maxillary first molars. Col2.3GFP was used as a marker to identify mature osteoblasts, cementoblasts and PDL fibroblasts. The effects of MTA after 2, 17, 30 days of injury, were examined and compared histologically to adhesive system sealing. The effects of two dilutions of medium conditioned with MTA on proliferation and differentiation of mesenchymal progenitor cells derived from bone marrow (BMSC) and periodontal ligament (PDL) *in vitro* were examined using presto blue viability assay, alkaline phosphatase and Von Kossa staining. The expression of markers of differentiation was assessed by real time PCR. Histological analyses showed better repair in teeth restored with MTA as shown by greater expansion of SMA9+ progenitor cells and Col2.3GFP+ osteoblasts compared to controls. The *in vitro* data showed that MTA conditioned medium reduced cell viability and osteogenic differentiation in both PDLs and BMSCs. In addition, cultures grown in the presence of MTA had marked decreases in SMA9+ and Col2.3GFP+ areas as compared to osteogenic medium confirming reduced osteogenesis. Thus, we concluded that MTA promotes regeneration of injured PDL and alveolar bone reflected as contribution of progenitors into osteoblasts. *In vitro*, MTA conditioned medium fails to promote osteogenic differentiation of both PDL and BMSC.

Descriptors: Periodontal Ligament; Stem Cells; Dental Materials.

Financial Support: This work has been supported by R01-AR055607 NIH/NIAMS to I.K and by R01-DE016689 & R90-DE022526 to M.M. BGM is supported by Connecticut Innovations grant 14-SCA-UHC. IOAQ is supported by fellowship from São Paulo Research Foundation (FAPESP)#2014/13750-0.