



## Z-003

### **Cytotoxicity analysis of different adhesive systems used in implant-retained maxillofacial prosthesis**

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#### **Objectives**

This study aimed to evaluate the cytotoxicity of different adhesive systems used to bond acrylic resin and facial silicone in implant-retained maxillofacial prosthesis, in keratinocytes cells (HaCat), through the analysis of cell proliferation, production of proinflammatory cytokines and extracellular matrix proteins

#### **Methods**

For this, 21 samples were divided into 7 groups: Resin, Silicone, Resin + Silastic Medical Adhesive Type A + Silicone (RAS), Resin + DC 1205 Primer + Silicone (RDCpS), Resin + Sofreliner Primer + Silicone (RSpS), Resin + DC1205 Primer + Silastic Medical Adhesive Type A + Silicone (RDCpAS), and Resin + Sofreliner Primer + Silastic Medical Adhesive Type A + Silicone (RSpAS). Eluates corresponding to 24 hours (h) of sample immersion in medium were prepared, in which HaCat cells were exposed for 72 h. Cytotoxicity was evaluated by cell viability MTT test. The Interleukin 1 $\beta$ , Interleukin 6 (IL6), Tumor Necrosis Factor  $\alpha$  and Macrophage Inflammatory Protein 1 $\alpha$  levels were evaluated by ELISA, and mRNA expression for Collagen Type IV (COL IV), Matrix Metalloproteinase 9 (MMP9) and Transforming Growth Factor  $\beta$  (TGF $\beta$ ), by RT-PCR. Data were submitted to ANOVA and Bonferroni tests ( $p < 0.05$ )

#### **Results**

The materials did not present cytotoxic potential. IL1 $\beta$ , TNF  $\alpha$  e CCL3/MIP1 $\alpha$  were not detectable. RDCpS presented the highest concentration of IL6. Concentrations of COL IV, MMP9 e TGF  $\beta$  were not statistically different among groups.

#### **Conclusions**

Materials evaluated were not toxic and DC 1205 primer not associated with adhesive (RDCpS group) may trigger an inflammatory process.

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