



0236

### **Gingival Keratinocyte Adhesion on Atomic Layer-Deposited Hydroxyapatite Coated Titanium**

F. A. Abushahba<sup>1, 2</sup>, S. Riivari<sup>1</sup>, N. Areid<sup>1</sup>, E. Närva<sup>3</sup>, E. Kylmäoja<sup>4</sup>, J. Holopainen<sup>5</sup>, M. Ritala<sup>5</sup>, J. Tuukkanen<sup>4</sup>, P. Vallittu<sup>2</sup>, T. Narhi<sup>1, 6</sup>

<sup>1</sup>Department of Prosthetic Dentistry and Stomatognathic Physiology, Institute of Dentistry, University of Turku, Turku, Finland, <sup>2</sup>Department of Biomaterials Science and Turku Clinical Biomaterial Center-TCBC, Institute of Dentistry, University of Turku, Turku, Finland, <sup>3</sup>Institute of Biomedicine and Cancer Research Laboratory FICAN West, University of Turku, Turku, Finland, University of Turku, Turku, Finland, <sup>4</sup>Department of Anatomy and Cell Biology, Research Unit of Translational Medicine, Medical Research Center, University of Oulu, Oulu, Finland, <sup>5</sup>Department of Chemistry, University of Helsinki, University of Helsinki, Helsinki, Finland, <sup>6</sup>Oral Health Care, Wellbeing services county of Southwest Finland, University of Turku, Turku, Finland

**Objectives** The aim of this study was to evaluate the effects of the atomic layer deposited hydroxyapatite (ALD-HA) coating on human gingival keratinocyte (HGK) adhesion, spreading, growth, and hemidesmosome formation on the titanium surface.

**Methods** Grade 2 square-shaped titanium substrates were prepared (n=62). The HA coating was done by first depositing with ALD CaCO<sub>3</sub>, which was hydrothermally converted to HA. Half the substrates were ALD-HA coated, while the other half was used as non-coated control (NC). The ALD-HA coating underwent surface characterization through scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDS) analysis. The initial cell adhesion and hemidesmosome formation of HGKs were evaluated after a 24-hour cultivation period. The cell proliferation was detected by cultivating cells for 1, 3 and 7 days. In addition, the levels of adhesion proteins integrin  $\alpha 6$  and  $\beta 4$  were detected with the Western Blot method. Furthermore, high resolution imaging of cell areas and adhesion protein signals was established using a confocal microscope.

**Results** SEM-EDS analysis demonstrated the formation of HA crystals on the ALD-HA surfaces. The relative cell attachment was significantly higher ( $p < 0.05$ ) on the ALD-HA surface compared to NC after 1 and 3 days of cell culture. No significant difference was found in integrin  $\alpha 6$  or  $\beta 4$  expression. The microscope evaluation showed significantly wider cells with peripheral hemidesmosome expression on ALD-HA surfaces compared to the NC ( $p = 0.0001$ ). The signal of laminin  $\gamma 2$  on the cell bottom layer was significantly higher on ALD-HA-coated surfaces compared to NC ( $p < 0.001$ ).

**Conclusions** Based on the findings of this in vitro study, the ALD-HA coating enhances the attachment of HGKs and promotes the expression of adhesion proteins on coated titanium surfaces. The results of this study indicate that ALD-HA coating has good potential for improving mucosal attachment on implant surfaces.