SCIENTIFIC GROUPS & NETWORKS: Implantology Research

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TITLE: The Influence of a Laser Modified Zirconia Implant Surface on Human Osteoblast In-vitro Behavior

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Objectives To evaluate the influence of laser passes in groove-texture zirconia implant surfaces on the in vitro response of human fetal osteoblasts

Methods Laser manufactured, meso-scale groove textured zirconia (YTZP) discs were produced using press-andsintering techniques. All surfaces were treated with Nd:YAG laser to produce 25μ m-spaced grooves. Each group had different number of laser passes (group A: 1; B: 2, C: 4 and D: 8 passes) in each groove site. Untextured zirconia discs were used as controls (UT). Sandblasted and acid-etched (SB-AE) protocol was applied to all samples to achieve 2.25 ± 0.42 µm mean surface micro-scale roughness. Human osteoblasts were cultured for 14 days by previously described methods. Cell morphology and adhesion were observed using scanning electron microscopy (SEM). Cell viability was evaluated at pre-defined time-points (1,3,7 and 14 days) using a commercial resazurin-based method. Alkaline phosphatase (ALP) activity of human osteoblasts was evaluated at 7 and 14 days using an enzymatic colorimetric assay. Collagen type I were evaluated at 3 days using enzyme-linked immunosorbent assay. All results were presented as mean ± standard deviation (SD). Group comparisons were tested using Anova (Tukey's post-hoc) using appropriate statistical software and significance was set at p<0.05.

Results Cell viability and proliferation increased over time for all groups, although without statistically significant differences between them (p>0.05), but significantly higher when compared to UT control (p<0.05) for 7 and 14 days evaluation. Collagen I levels were higher for all groups when compared to UT control (p<0.05) and ALP activity was significantly increased in group D and UT control when compared to other groups at 14 days (p<0.05). **Conclusions** Osteoblast viability, proliferation and differentiation were significantly enhanced by laser surface groove texturing. A tendency towards an enhanced osteoblast differentiation for higher number of laser passes should be further investigated.