

World Congress on Joint Event On
**NOVEL TRENDS AND ADVANCES IN BIOTECHNOLOGY,
CELL & STEM CELL RESEARCH &**
15TH ANNUAL CONGRESS ON PEDIATRICS
November 28-29, 2018 Barcelona, Spain

Structural and biochemical analysis of extracellular matrix formed by jaw periosteal cells

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Statement of the Problem: For clinical applications of stem cell-based therapies, animal component-free culture conditions are required. In previous studies, we detected robust mineralization of Jaw Periosteal Cells (JPCs) under cultivation with clinical approved Human Platelet Lysate (hPL) compared to FCS conditions. In the present study, we performed qualitative analyses of the Extracellular Matrix (ECM) formed by JPCs under hPL and FCS culture conditions.

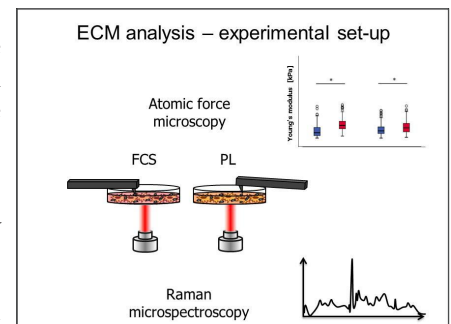
Methodology & Theoretical Orientation: JPCs were induced osteogenically under FCS and hPL culture conditions. At the end of osteogenic differentiation, biochemical composition of the formed precipitates was analyzed by Raman microspectroscopy and mechanical properties were assessed by Atomic Force Microscopic (AFM) measurements.

Findings: By Raman analyses, higher phosphate, lower carbonate content and higher crystallinity of hydroxyapatite minerals were detected under hPL culture conditions. Furthermore, regarding the quality of the collagen network, higher ratios of proline/hydroxyproline and higher collagen cross-linking were detected in hPL-cultured JPCs. Since hPL-supplementation leads to nearly equal production of the precursor protein proline and the mature protein hydroxyproline, these findings might indicate higher elastic properties of the collagen network in contrast to the FCS supplementation. However, cross-linking seems to be higher under hPL culture conditions. When areas of cell monolayers without precipitates were monitored by AFM, higher Young's modulus of untreated and osteogenically induced JPCs under FCS compared to hPL culture conditions were detected. The opposite was the case for the formed precipitates: under hPL conditions, precipitates formed by JPCs showed significantly higher Young's modulus than those formed under FCS culturing.

Conclusion & Significance: In the present study, we detected significant differences in the biochemical composition and the mechanical properties of the mineralized extracellular matrix formed by hPL and FCS-cultured JPCs. The combined technologies represent an optimal tool to assess the quality of *in vitro* formed bone tissue.

Biography

Alexander has her expertise in bone tissue engineering. The aim of her research focus is to develop an optimal stem cell source and appropriate biomaterials for future regeneration of maxillofacial bone defects. Therefore, her research group characterizes jaw periosteal cells and their *in vitro* osteogenic capacity and establishes optimal culture conditions and biofunctionalization strategies for biomaterials in view of future clinical performance. Related thereto, the establishment of suitable cell imaging approaches in order to assess cellular dynamics and biochemical changes of living cells represents a present focus of Alexander's work group.



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