Three dimensional visualization of engineered bone and soft tissue by combined x-ray micro-diffraction and phase contrast tomography

<u>Alessia Cedola^{1*}</u>, Gaetano Campi^{2*}, Daniele Pelliccia³, Inna Bukreeva¹, Michela Fratini⁴, Manfred Burghammer⁵, Luigi Rigon^{6,7}, Fulvia. Arfelli^{6,7}, Giuliana Tromba⁸, Maddalena Mastrogiacomo⁹

¹Istituto per i Processi Chimico Fisici - CNR, c/o Physics Dep. at 'Sapienza' University, Rome-ITALY. ²Istituto di Cristallografia, CNR, via Salaria Km 29.300, 00015 Monterotondo Roma, Italy.

⁵ESRF- Grenoble FRANCE

⁶Dipartimento di Fisica, Università di Trieste, 34127 Trieste, Italy

⁷Istituto Nazionale di Fisica Nucleare, Sezione di Trieste, 34127 Trieste, Italy

⁸Sincrotrone Trieste SCpA, 34149 Basovizza (Trieste), Italy

⁹Dipartimento di Medicina Sperimentale dell'Università di Genova & AUO San Martino-IST Istituto Nazionale per la Ricerca sul Cancro, Largo R. Benzi 10, 16132, Genova, Italy

*Alessia Cedola, E-mail address: alessia.cedola@cnr.it

Computed X-ray phase contrast micro-tomography is the most valuable tool for a threedimensional and non destructive analysis of the tissue engineered bone morphology. We used a Talbot interferometer for a precise 3D reconstruction of both bone and soft connective tissue, regenerated in vivo within a porous scaffold. For the first time the X-ray tomographic reconstructions have been combined with X-ray scanning micro-diffraction measurement on the same sample, in order to give an exhaustive view of the role of the different tissues participating to the biomineralization process. In particular our experimental approach allows for a deeper understanding of the role of collagen matrix in the organicmineral transition, which is a crucial issue for the development of new bio-inspired composites.



Figure 1. a) Diffracted intensity, collected in different positions, indicating three different tissues, namely soft tissue (ST), due mainly to the collagen fibres, scaffold (SC) and newly formed bone tissue (B). From integration we obtained the spatial distribution of the bone, collagen and ACP, shown in figure 1b, 1c, 1d respectively. Figure 1e is the δ (real part of the refraction index) distribution image obtained by the tomographic reconstruction.

³School of Physics, Monash University, Victoria 3800, Australia

⁴Centro Fermi, compendio Viminale, Piazza del Viminale 1 - Roma - 00184 - Italy