

Monitoring bioinspired collagen mineralization through X-Ray Total Scattering

JOSÉ MANUEL DELGADO-LÓPEZ^{1,2}, FEDERICA BERTOLOTTI^{1,3,4}, JAN SKOV PEDERSEN³, ANTONIO CERVELLINO⁵, NORBERTO MASCIOCCHI^{1,4} AND ANTONIETTA GUAGLIARDI^{1,6}

¹Total Scattering Laboratory (To.Sca.Lab), Como, Italy.

²IACT (CSIC-UGR), Granada, Spain.

³iNANO, Aarhus University, Denmark.

⁴DiSAT, Università dell'Insubria, Como, Italy.

⁵Swiss Light Source, PSI, Switzerland.

⁶Istituto di Cristallografia, IC-CNR, Italy.

Bone is an extremely complex tissue with many levels of organization.^[1] The building blocks of such a complex architecture are the mineralized collagen fibrils.^[1] Over the past years, notable advances have been made towards unveiling the role of the organic matrix (*i.e.*, collagen fibrils and non-collagenous proteins, NCPs) in driving mineral nucleation and growth. However, many aspects are still far to be understood. For instance, the mechanisms underlying the amorphous-to-apatite crystallization and clarifying the origin of platy apatite crystals has never been directly revealed in neither *in vivo* nor *in vitro* experiments. Remarkably, spherical amorphous calcium phosphate (ACP) precursors are found at the early stages, whereas apatite platelets are observed as the final crystalline product.^[2]

Results from the *in situ* monitoring of collagen mineralization (under physiological conditions)^[3] by synchrotron-based SAXS and WAXTS will be discussed. The great potential of Total Scattering methods (TS) for providing the stoichiometry, the structure as well as size and shape distributions of nano-apatites^[4] will be also discussed.

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References

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