

ICVS/3B's



# 3<sup>rd</sup> 3B's Symposium on Biomaterials and Stem Cells in Regenerative Medicine

3B's Research Group Auditorium– AvePark, Caldas das Taipas, Guimarães, Portugal Date: 22 May, 2013



**Chairmen:** João F. Mano and Rui L. Reis 3B's Research Group, University of Minho, Portugal





# Program

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Program		
9:00-09:10	Welcome	João F. Mano / Rui L. Reis
	Stem cells and biological aspects in regenerative medicine (Chair: Rui L. Reis)	
9:10-9:35	Adult stem cells for TERM: sources and manipulation	Ana Rita Costa-Pinto
9:35-10:00	Induced pluripotent stem cells	Ana M. Martins
10:00-10:25	Vascularization strategies in Regenerative Medicine	Rogério R. Pirraco
10:25-11:00	Keynote Lecture: Extracellular matrix-driven tissue regeneration BREAK	Alexandra P. Marques
	Biomaterials: from <i>nano</i> to <i>macro</i> (Chair: Ricardo A. Pires)	
11:20-11:45	Polymeric biomaterials from marine-origin	Anabela Alves
11:45-12:10	What is common between pintarolas and cells?	Iva Pashkuleva
12:10-12:35	Nanobiomaterials in tissue engineering LUNCH BREAK	Albino Martins
14:00-14:25	Shaping biomaterials into porous 3D constructs	Ana R. Duarte
14:25-14:50	Shaping biomaterials into spherical objects	Clara R. Correia
14:50-15:15	Combinatorial analysis of biomaterials for Tissue Engineering	Mariana B. Oliveira
	Using natural-based biomaterials in case studies of Tissue Engineering	
	(Chair: João F. Mano)	
15:40-16:00	System	Susana R. Cerqueira
16:00-16:20	Skin tissue engineering	Mariana Cerqueira
16:20-16:40	Cartilage tissue engineering	Marta L. Silva
16:40-17:00	Regeneration of the intervertebral disk	Joana S. Correia
17:00-17:20	Strategies for the regeneration of the tendon	Márcia Rodrigues
17:20-17:40	Sports & regenerative medicine	Hélder Pereira
	Creative thinking	
	(Chair: Nuno M. Neves)	
17:40-18:15	Keynote Lecture: 3C's - Cultura, Ciência e Criatividade em gastronomia	Renato Cunha





## TISSUE ENGINEERING AS A REMARKABLE TOOL FOR CARTILAGE REPAIR

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Articular cartilage is a very specialized tissue with outstanding load-bearing capacity. It consists mainly of a dense extracellular matrix (ECM) with chondrocytes embedded on it. Cartilage has very low capacity of self-repair and regeneration after traumatic, degenerative or inflammatory injury. Current available surgical treatments for cartilage repair present several drawbacks, such as possible implant rejection or infection, or the need for revision after some years of implantation. Autologous chondrocyte implantation (ACI) is an autologous therapy that was proposed as a basis for tissue engineering strategies to repair cartilage (1). Modifications on various aspects of this surgical technique have been developed, comprising the use of natural-based scaffolds as supports for chondrocyte expansion (2).

Many strategies and systems have been developed along the years for cartilage regeneration and repair. Scaffolds play a major role in those strategies, as they provide the support for cell growth and to promote extracellular matrix production. Both natural based (3) or synthetic scaffolds (4) have been successfully used as supports for chondrogenic differentiation or cartilage-like tissue production.

The interest in cells cross-talk and communication has been growing in the past years, revealing that signalling pathways are pivotal elements when understanding the tissue formation and its repair mechanisms (5). Chondrocytes release morphogenetic signals that influence the surrounding cells, for example, stem cells, to differentiate into the chondrogenic lineage (5). In fact, the increased cartilage formation on co-cultures using stem cells and articular chondrocytes has been reported (6). Therefore, the study of co-cultures using chondrocytes and undifferentiated cells is a very promising strategy to develop engineered cartilage.





## References

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Welsch GH, Trattnig S, Hughes T, Quirbach S, Olk A, Blanke M, et al. T2 and T2\* mapping in patients after matrix-associated autologous chondrocyte transplantation:

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6. Meretoja VV, Dahlin RL, Kasper FK, Mikos AG. Enhanced chondrogenesis in cocultures with articular chondrocytes and mesenchymal stem cells. Biomaterials. 2012 Sep;33(27):6362-9.