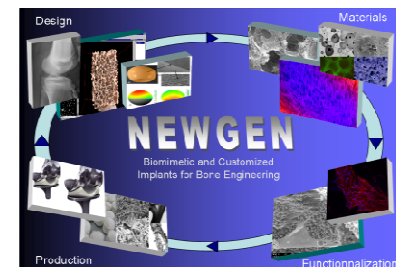


- ✓ **Complete denomination:** Histocell S.L.
- ✓ **Location (city, country):** Derio, Spain
- ✓ **Director:** Dr. Julio Font
- ✓ **Contact person in NEWGEN:** Dr. Begoña Castro
- ✓ **Working Group involvement:** WG 1, WG3 & WG4
- ✓ **Staff:** 24 staff, (12 of them PhD), fully dedicated to research and manufacturing activities.
- ✓ **Research topics:** Bone regeneration, Advanced wound healing, Inflammatory diseases, Neuroregeneration, Ischemia and Lung therapy.
- ✓ **Researchers expertises:** Development of new methods and technologies to apply them in regenerative medicine, and uses Adult Mesenchymal Stem Cells (AMSCs) and Bioactive materials (scaffolds) as key technology.



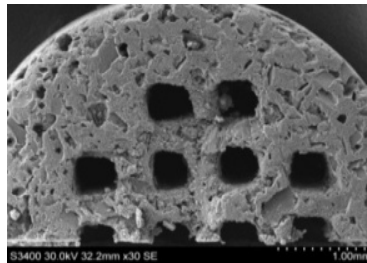
Histocell

Parque Tecnológico de Bizkaia
48160, Derio - SPAIN



COST Action MP1301

SYNTHETIC BONE SUBSTITUTE FOR OSSEOUS REGENERATION



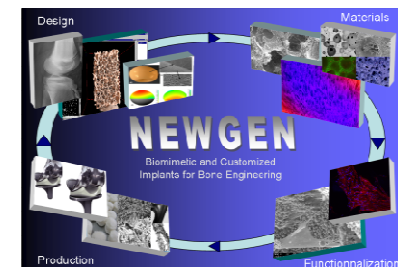
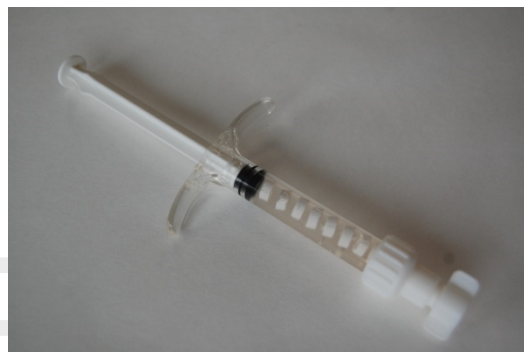
Description: Resorbable cylinder-shaped matrixes composed of monetite featuring a combined micro/macroporous structure

Target: To be directly implanted into bone defects caused mainly by traumatic injuries, pseudoarthrosis, tumor resection surgery or osteomyelitis



MA: Conducts a guided bone remodelling process. Enables osteosynthesis and angiogenesis. New tissue formation occurs parallel to absorption

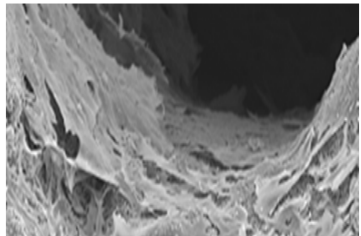
IP status: PCT/ES2009000358 (2009)/Patent granted in Mexico
Development: *Preclinical studies ongoing*



COST Action MP1301



BONE REGENERATION



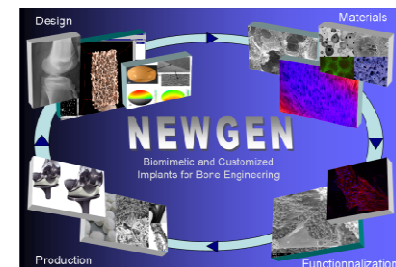
Product: Autologous adipose tissue derived MSCs grown in an in house developed, engineered bone substitute with optimal bone remodeling properties

Target: Severe bone defects caused by pseudoarthrosis, surgery or traumatic injuries

Innovation: Regenerative capabilities of cells are supported by the use of a bone-like scaffold made of a biocompatible calcium phosphate derivate. The scaffold's nature enables a directed osseous remodeling process for an optimal integration of the newly formed tissue with the surrounding bone and ensures its adequate revascularization

Development stage: Preclinical phase completed successfully. Phase I trial scheduled for early 2015

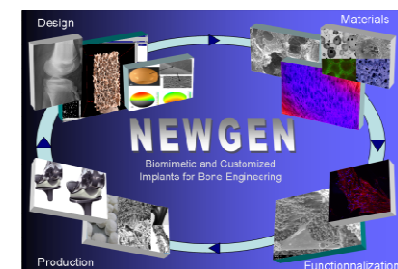
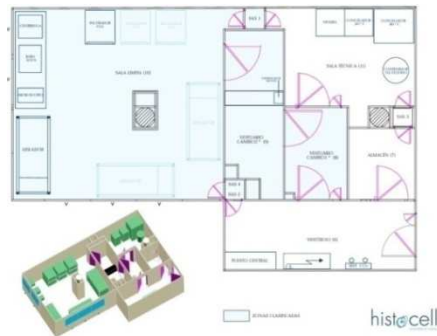
Status: Licensing & codevelopment agreement with Salvat



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R&D Area

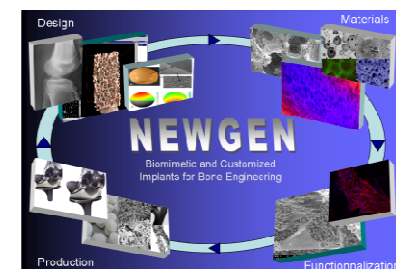
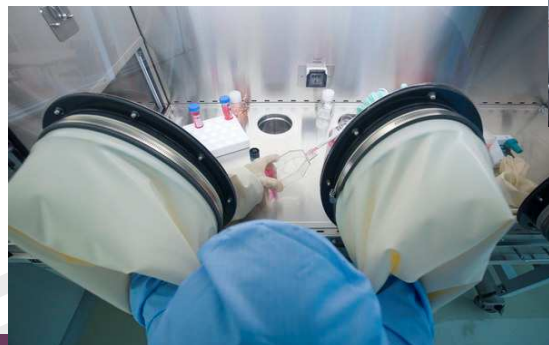
567 m2 working area composed by completely equipped offices and laboratories: Molecular Biology Laboratory, Cell Culture Laboratory, Cytofluorometry, Spectrophotometry, Histology Immunomarking...



GMP Manufacturing facility

65 m2 authorized clean room

GMP compliance certification from the Spanish drug and medical device agency (AEMPS) for the manufacturing of clinical grade Mesenchymal Stem Cells and Chondrocytes.



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