



UNIVERSIDADE DA BEIRA INTERIOR
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Electrospun poly(ϵ -caprolactone) nanofibers for bone regeneration and other biomedical applications

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Biomedical Sciences
(2nd cycle of studies)**

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**Produção de nanofibras de poli(ϵ -caprolactona)
por electrospinning para futura aplicação na
regeneração óssea e outras aplicações biomédicas**

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Dissertação para a obtenção do Grau de Mestre em
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“Imagination is more important than knowledge. For while knowledge defines all we currently know and understand, imagination points to all we might yet discover and create.”

Albert Einstein

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Abstract

Bone tissue is a complex and hierarchical structure with many functions in the body. Although this tissue has the capability for self-generation, large bone defects due to various diseases or fractures may need clinical treatment. However, the current clinical treatments are based on bone grafts and other bone substitutes, which have several limitations. Tissue engineering is a multidisciplinary field that emerged from the need to extinguish these clinical limitations. This vast field of science uses various tools in seeking for effective tissue regeneration. In this context, this work aimed the production of functional materials that mimic the nanostructure of bone tissue and therefore the cellular microenvironment, promoting bone regeneration. In this way, an electrospinning apparatus was mounted and optimized for the production of polycaprolactone nanofibers. Additionally, several electrospinning parameters that influence the morphology of the electrospun nanofibers produced were studied. For application in bone regeneration, the combination of polycaprolactone nanofibers with β -tricalcium phosphate scaffolds, mechanically more resistant, was investigated. Moreover, a preliminary study about the capacity for this system to allow controlled release of biomolecules was conducted through the incorporation of a model protein into the nanofibers. In order to characterize the biological properties of the systems produced, *in vitro* cytotoxicity assays were performed. These assays revealed that the polycaprolactone nanofibers produced are biocompatible and that the coating of β -tricalcium phosphate scaffolds with these nanofibers improve this biological performance, when compared to ceramic scaffolds without coating. The potentiality of nanofibers herein produced was also evaluated for the modification of microfiltration membranes. These membranes showed a large increase in the plasmid DNA rejection.

Keywords

Electrospinning, polycaprolactone, nanofibers, tissue engineering, coating of scaffolds, modification of membranes.

Resumo

O tecido ósseo é uma estrutura complexa com diversas funções no organismo. Apesar deste tecido possuir uma capacidade de auto-regeneração única, defeitos ósseos com uma grande extensão causados por doenças ou fracturas podem necessitar de tratamento em meio hospitalar. No entanto, estes tratamentos são algo limitados, pois baseiam-se em transplantes e em substitutos ósseos compostos por materiais inertes. A engenharia de tecidos é uma área multidisciplinar que emergiu da necessidade de extinguir estas adversidades. Este vasto campo científico utiliza diversas ferramentas na procura de uma regeneração de tecidos mais eficaz. Neste âmbito, este trabalho pretende produzir materiais funcionais que mimetizem a nanoestrutura do tecido ósseo e, portanto, o microambiente celular, favorecendo a regeneração óssea. Com base neste pressuposto, montou-se um sistema de electrospinning e procedeu-se à sua optimização para a produção de nanofibras de policaprolactona. Adicionalmente, diversos parâmetros que influenciam a morfologia das nanofibras produzidas por electrospinning foram estudados. Para aplicação na regeneração óssea, a combinação das nanofibras de policaprolactona com scaffolds de β -tricálcio fosfato, mais resistentes mecanicamente, foi investigada. Adicionalmente, foi realizado um estudo preliminar sobre a capacidade deste sistema permitir a libertação controlada de biomoléculas, através da incorporação de uma proteína modelo nas nanofibras. O perfil citotóxico dos sistemas produzidos foi caracterizado através de ensaios *in vitro*. Estes estudos revelaram que as nanofibras de policaprolactona produzidas são biocompatíveis e que o revestimento dos scaffolds de β -tricálcio fosfato com estas nanofibras melhora as propriedades biológicas em relação aos scaffolds cerâmicos sem revestimento. A potencialidade das nanofibras produzidas foi ainda testada na modificação de membranas de microfiltração. Este estudo demonstrou um aumento na rejeição de DNA plasmídico após o revestimento da membrana com as nanofibras.

Palavras-chave

Electrospinning, policaprolactona, nanofibras, engenharia de tecidos, revestimento de scaffolds, modificação de membranas.

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List of Acronyms

ALP	Alkaline phosphatase
BMPs	Bone morphogenetic proteins
BSA	Bovine serum albumin
DMEM-F12	Dulbecco's modified eagle's medium
EDTA	Ethylenediaminetetraacetic acid
EGF	Epidermal growth factor
FBS	Fetal bovine serum
FDA	Food and Drugs Administration
HA	Hydroxyapatite
RO	Reverse osmosis
MF	Microfiltration
MSCs	Mesenchymal stem cells
MTS	3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium reagent, inner salt
NF	Nanofiltration
PCL	Poly(ϵ -caprolactone)
pDNA	Plasmid DNA
SEM	Scanning electron microscopy
TCP	β -Tricalcium phosphate
TGF- β	Transforming growth factor - β
UF	Ultrafiltration
VEGF	Vascular endothelial growth factor
3D	Three-dimensional

Chapter I

Introduction

1. Introduction

1.1 Bone tissue

1.1.1 Formation and composition of bone

As part of the skeletal, bone plays several critical functions in the body: it is involved in structural support, is fundamental for locomotion, ensures protection for the internal organs (chest, cranium, spine), provides tissue base for the production of blood cells (haematopoiesis) and immune cells and form a mineral reservoir to maintain electrolyte homeostasis in the body [1, 2].

Bones are formed through two distinct pathways (figure 1): endochondral ossification gives rise to long bones, which include appendicular skeleton, facial bones, vertebrae and lateral clavicles; intramembranous ossification gives rise to the flat bones, comprising the cranium and medial clavicles [3]. Both types of ossification involve an initial condensation of mesenchyme, where cells are positioned adjacent one to another, and the successive formation of calcified bone [4]. However, intramembranous ossification accomplishes this directly, whereas endochondral ossification has an intermediate step in which an avascular tissue (cartilage) is formed before being mineralized and converted into bone tissue [3, 5].

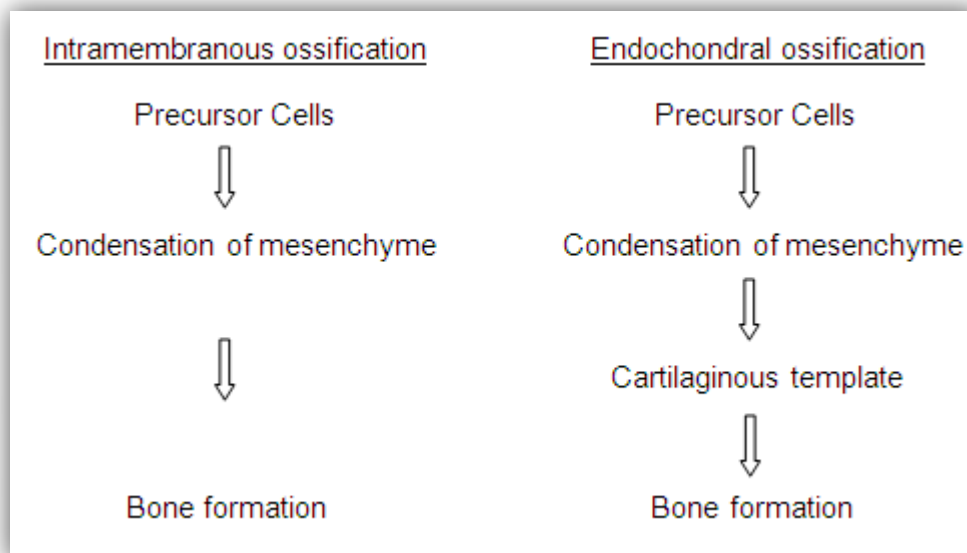


Figure 1 - Bone developmental processes (Adapted from [3]).

Bone tissue is composed by mineral components (70%) and organic components (30%). The first one is almost all composed of an analogous form of hydroxyapatite [$\text{Ca}_{10}(\text{PO}_4)_6(\text{OH}_2)$], that can incorporate ions like magnesium, zinc, sodium, potassium, chlorine, fluorine and

strontium [6]. The high surface/volume ratio and the crystals irregularity make them more soluble and easily mobilized to metabolic demands [6].

The organic part of the bone is composed by 2% of bone cells and 98% of organic components from the extracellular matrix (ECM) [6]. The principal organic component of ECM is collagen type I. This protein is a heteropolymer of three chains, each of them having the primary structure (Gly-X-Y)_n, where X and Y are frequently proline or hydroxyproline [7]. Collagen is deposited in matrix along the mechanical stress lines and provides a backbone for the deposition of bone mineral [3, 6]. The mechanical properties of a bone are determined by this combination of materials. The inorganic phase is related to the resistance during compression, whereas the organic phase is responsible mainly for the viscoelastic properties in tension or shear [8].

The cellular elements in bone act as metabolic regulators and are responsible for the continuous remodelling of bone tissue [6]. Three types of cells coexist in bone: Osteoblasts are derived from marrow stromal fibroblastic stem cells. These cells are involved in the synthesis of a decalcified organic matrix, mainly constituted by collagen type 1, called osteoid. In this process, they produce large amounts of alkaline phosphatase, an enzyme that prepares the matrix for mineralization [9]; Osteocytes (derived from osteoblasts) are surrounded during the mineralization process in bone lacunae that have a network of canaculi, through which they make their metabolic exchanges and regulate the blood-calcium homeostasis [3, 6]; Osteoclasts (derived from the monocyte-macrophage lineage) are giant cells that produce enzymes, like acidic phosphatase, and are responsible for bone resorption [10].

Bone tissue is maintained by a balance of bone-forming and bone-resorbing cells. A malfunction in this equilibrium results in an excessive bone resorption that leads to anomalous conditions including osteoporosis, osteoarthritis, periodontitis and metastatic bone disease [11, 12].

1.1.2 Structure of bone

Morphologically bone can be described based on its density: cortical (compact) bone and cancellous (spongy or trabecular) bone [3, 11].

Cortical bone has as fundamental unit the osteon or Haversian system. Osteons consist of a central canal (Haversian canal) derived from intraosseous arteries and perforating canals surrounded by concentric rings of the matrix [11]. Due to the preferential orientation of osteons along the direction of the forces supported by the diaphysis, that arteriole is almost always disposed to length of bone. The Haversian canal communicates to each other by horizontal branches called Volkmann canals [6].

Cancellous bone presents a structure of intersecting trabeculae, following mechanical force lines, giving them a greater resistance to compressive forces. On the other hand,

cortical bone is better able to withstand forces of extension-flexion and torsion forces [6]. The network of trabeculae is covered by an epithelial membrane of osteoblasts that can be activated to form more bone lamellae. On the trabecular surface the osteoclasts are involved in resorbing bone matrix and releasing minerals in Howship lacunae [6].

A schematic representation of both bone types can be observed in figure 2.

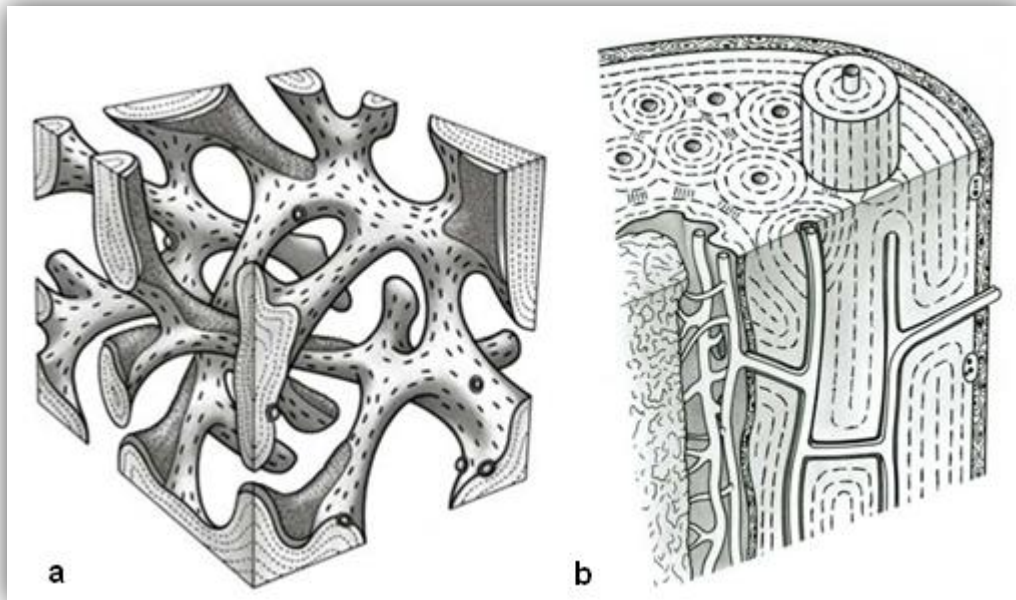


Figure 2 - Structure of bone: a) Cancellous bone. b) Cortical bone (Adapted from [3]).

The difference between cancellous and cortical bone is not only structural but also functional: cortical bone provides mechanical and protective functions, while cancellous bone is involved in metabolic functions, like the calcium homeostasis [3].

The hierarchical arrangement of bone tissue is represented in figure 3.

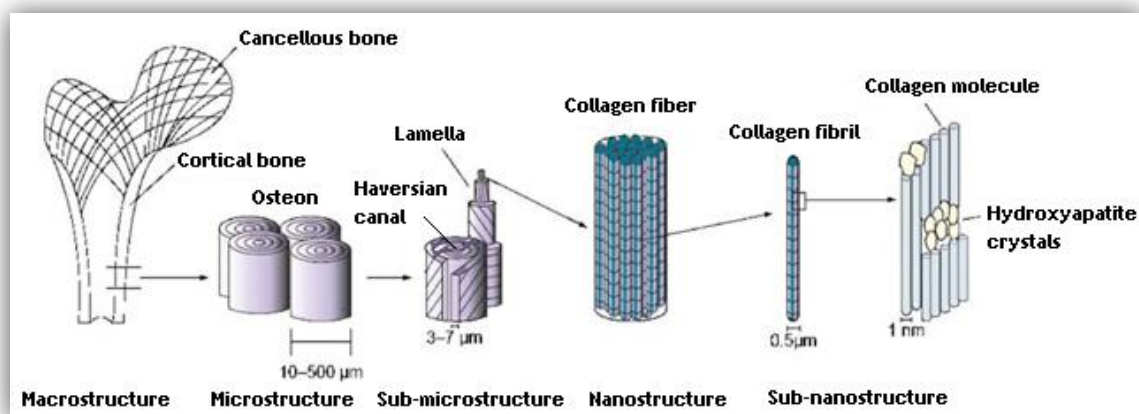


Figure 3 - Illustration of the macro to nano structures of the bone (Adapted from [13]).

1.1.3 Bone self-regenerative ability

Bone is highly vascularized and a very dynamic tissue, since it has both high mechanical properties and regenerative capacity [14]. This regenerative property is crucial for skeletal homeostasis and fracture repair. Unlike soft tissue healing, the regeneration of the natural bone tissue occurs without scar formation [15]. Fracture healing can be divided into three phases that form a continuous healing process after injury. In temporal order, this process comprises an inflammatory phase, a reparative phase and a remodeling phase [15, 16]. The overall process of bone repair is illustrated in figure 4.

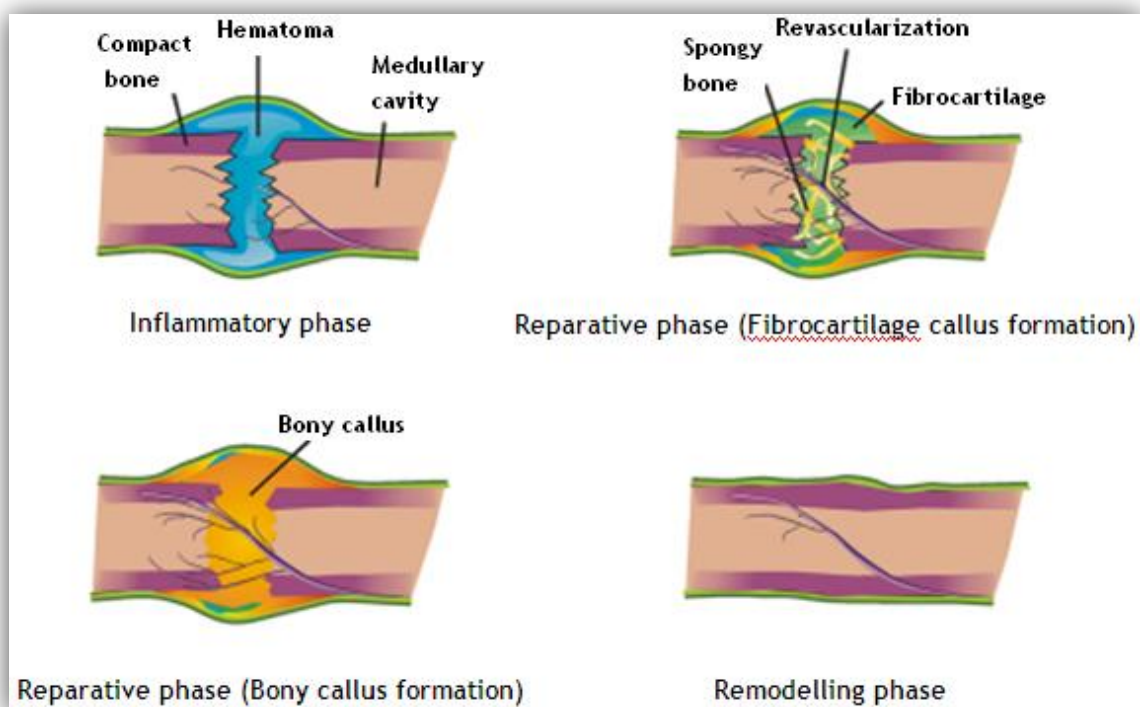


Figure 4 - Schematic representation of fracture healing (Adapted from [15]).

Inflammatory phase

At the onset of bone damage, an inflammatory response is elicited during 1 week post fracture. This reaction helps to immobilize the fracture in two ways: pain causes the individual to protect the injury and the enlargement of the site (swelling) keeps the fracture from moving. The blood vessels rupture results in the activation of the complement cascade, platelet aggregation, and release of its granule contents, which triggers chemotactic signals [15, 16]. Hemostasis is guaranteed by platelets and by the formation of a hematoma. On the other hand, the hematoma become a fibrin network that provides pathways for cellular migration and serves as a source of signal molecules, that initiate cellular events essential to

fracture healing. Lymphocytes, blood monocytes, macrophages and osteoclasts are attracted to the injury site and are activated to “clean up” the wound of biological debris and to release cytokines that can stimulate angiogenesis [15, 16]. Growth factors that are released include insulin-like growth factor, vascular endothelial growth factor (VEGF), platelet-derived growth factor, transforming growth factor- β (TGF- β) and epidermal growth factor. These growth factors play important roles in the proliferation and differentiation of the mesenchymal stem cells (MSCs) [16].

Reparative phase

The reparative phase occurs before the inflammatory phase subsides and lasts for several weeks. A reparative callus tissue is formed at fracture site, which will be eventually replaced by bone. This callus enhances mechanical stability of the site by supporting it laterally. At this phase, the callus comprises of fibrous connective tissue, blood vessels, cartilage, woven bone and osteoid. While fracture wastes are being resorbed, pluripotent MSCs begin to differentiate in fibroblasts, chondroblasts and osteoblasts. As bone healing proceeds, the pH gradually becomes neutral and then slightly alkaline, which is optimal for alkaline phosphatase (ALP) activity and its role in the mineralization of the callus. The reparative phase includes endochondral bone formation and intramembranous bone formation [15, 16]. As the mineralization process evolves, the callus calcifies becoming more rigid and the fracture site is considered internally immobilized. During this stage, collagen fibers are not perfectly aligned and the callus is composed of an unorganized woven bone, which connects the two fracture ends. Proteins that regulate this phase include bone morphogenetic proteins (BMPs), parathyroid hormone-related peptide, osteopontin, osteocalcin, ALP and bone sialoprotein [16].

Remodeling phase

The remodeling phase is the final phase in fracture healing. In this phase the unorganized bone woven is replaced with a more organized structure and the excess of callus is resorbed. Remodeling of fracture repair consists of osteoclastic resorption of poorly located trabeculae and formation of new bone along lines of stress. This results in a gradual modification of the fracture region under the influence of mechanical loads, until an optimal stability is achieved and the bone tissue acquires similar architecture to that it had before the fracture occurred [15, 16]. The remodeling process is guided by expression of specific genes, protein synthesis and secretion. Osteoblast-osteoclast homeostasis is balanced by factors like macrophage colony-stimulating factor, tumor necrosis factor- α , receptor activator of nuclear factor κ B and osteoprotegerin [16]. The events of this stage continue throughout

adult life. Changes in estrogen levels, such as those occurring in postmenopausal women, can alter bone homeostasis; hence the bone loses its natural stability, what explains the increased susceptibility of women in comparison with men to bone fractures with aging [16].

1.1.4 Clinical treatment of bone: motives and purposes

In general, fracture healing is completed after 6-8 weeks after the initial injury occurs [15]. However, the healing capacity of bone has its limitations. Large bone defects due to trauma, fractures, cancer, periodontitis, osteoporosis and infectious diseases may not heal by themselves and result in non-union [17-19]. The size of bone fractures and defects that can be self-regenerated is designated as the “critical size defect” and will not heal during the lifetime of the patient. Thus, for large defects, medical intervention is often required in order to allow bone regeneration [14, 20, 21].

The intervention mentioned above, requires the introduction of implantable materials as bone grafts to promote the healing process [22]. Autografts (bone transplanted from the self-patient) are considered the gold standard for bone repair due to their osteoconductivity, osteoinductivity and biocompatibility without the risk of pathogen transfer [17]. Although autografts exhibit the best clinical outcome, they suffer from several disadvantages such the limited quantity, size and shape of donor tissue, and the requirement of surgery at multiple sites to harvest donor tissue from the patient [46-49]. Allografts (bone transplanted from one individual to another of the same species) are osteoconductive and more abundant in supply than autografts, but carry out the risk of pathogen transmission and immune response, which can lead to graft rejection [17, 23, 24]. Xenografts (bone transplanted from animals of different species) hold risks of immune response and pathogen transfer between species, thus limiting their use [1, 21]. As a result, there is a constant need for bone substitutes due to severe bone injuries, degenerative diseases and reconstructive surgery. In recent times, there was a remarkable increase in the use of biomaterials for bone related surgical applications [25]. However, a gap remains to be filled since no synthetic material used until now, presents the native properties of bone, attending both biological aspects as well as mechanical requirements [25].

1.2 Tissue engineering

In the last decade of the past century, tissue engineering emerges as a perspective to abolish the limitations of tissue grafts [26]. Tissue engineering is defined as an interdisciplinary field of research that applies methods of materials engineering and life sciences to create artificial constructs that restore, maintain or enhance tissues and organs functions [27, 28]. This field include a wide range of strategies employing cells, scaffolds, cytokines and genetic manipulation for the regeneration of tissue *in vivo* or the production of tissue *in vitro* [29]. Scaffolds are exogenous extracellular matrices designed to allow cell adhesion and proliferation and also to provide mechanical support until the newly formed tissue is structurally stabilized [30]. One approach to design exogenous extracellular matrices for tissue engineering is to mimic the functions of the extracellular matrix (ECM) components naturally found in tissues [30]. All artificial extracellular matrices used to engineer tissues have three fundamental roles: first, they facilitate the localization and delivery of cells to specific sites in the body; second, they define and maintain a structure for the formation of new tissues; and finally they guide the development of new tissues with their appropriate functions [30].

In bone tissue regeneration scaffolds are usually biodegradable and act as a temporary matrix for cellular proliferation and for extracellular matrix deposition. In addition, scaffolds work as a model for vascularization of new tissue (angiogenesis) and can also serve as carriers for biologically active agents, like growth factors that can enhance the regenerating potential of the system [31, 32].

An important aspect of tissue engineering science is the investigation of the interactions of cells with absorbable matrices and bioactive molecules (growth factors and cytokines), involved in the formation of new tissue [29]. The cells behaviour is strongly determined by a complex interaction of physical and chemical signals at the nanoscale [33]. Therefore, the incorporation of biological factors in scaffolds is common practice for tissue engineering applications. The purpose is to create a bioinspired extracellular environment that directs specific cell responses through activation of signalling cascades. These cascades will modulate gene expression and control important cell processes to allow tissue regeneration [33].

1.2.1 Biomaterials for bone tissue engineering

A biomaterial has been defined in the literature as “a substance that has been engineered to adopt a structure which, alone or as part of a complex system, is used to direct the course of any therapeutic or diagnostic procedure, in human or veterinary medicine” [34]. The biomaterials are used in tissue engineering for the manufacture of scaffolds that

provide structural support for specific cell attachment, spreading, migration, proliferation and differentiation [22, 31].

The first approach in the choice of a material to be used in the manufacture of a scaffold should be the evaluation of *in vitro* cytotoxicity, by using cell culture. Thus, the selection of the material to produce a scaffold is a crucial step in the construction of a tissue engineered product, since its properties are fundamental for the successful application of this three-dimensional (3D) matrix in the production of biomedical devices [32].

Ideally, scaffolds for bone tissue engineering should be biocompatible for a good integration into host tissue without eliciting an immune response; osteoconductive, i.e. should possess interconnected pores with a proper size to allow cell infiltration, neovascularization and removal of metabolic wastes; should be mechanically resistant to withstand local stress and provide structural support during the new bone tissue growth and remodelling [17, 35, 36]. The degradation products of the biomaterial should not be toxic and should be safely removed from the biological system. If this degradation rate is finely tuned, the scaffold is degraded in such a way that the scaffold is completely resorbed by the time that the bone defect is totally regenerated [17]. In addition, scaffolds should also possess a bioactive behaviour, promoting osteointegration and stimulating the growth and differentiation of bone cells, as well as be able to be sterilized without losing their bioactivity [25, 37].

A vast diversity of materials including ceramics, composites and polymers has been studied for the production of scaffolds for bone tissue engineering. These materials can be synthetic or natural, have different properties and show different degradation rates [22, 37, 38].

Ceramics such as hydroxyapatite (HA), calcium phosphates and bioactive glasses have been attracted special attention, due to their excellent biocompatibility along with their osteoconductive and osteoinductive properties [39]. Depending on their composition, particle size and production process, ceramics can have various degrees of bioactivity, which is the ability to chemically bond and be integrated into living bone through the formation of HA [40]. However, these materials are brittle and have low mechanical stability, making them unsuitable for load bearing applications [41].

Biodegradable polymers can avoid some of the drawbacks of ceramics. Natural polymers obtained from animal or vegetal sources, include proteins such as collagen, fibronectin, and polysaccharides such as chitosan, starch and hyaluronic acid [22, 42]. The major advantages of these materials are their great structural similarity to natural ECM constituents, their low immunogenic potential, as well as their high abundance [22, 42]. Synthetic polymers have great advantages because their structure, composition and, consequently, their properties can be tailored to specific needs [22, 43]. Polymers biodegradability can be controlled by different ways. Some polymers can suffer hydrolysis, others can be degraded by enzymatic pathways [43]. In addition, they are more ductile than ceramics and some can, in their solid form, reach mechanical compression strength close to

that of cortical bone [43]. The most widely investigated are poly(lactic acid), poly(glycolic acid), poly(ϵ -caprolactone) (PCL) and their copolymers such as poly (L-lactic-co-glycolic acid) [44-46]. Among these polymers, PCL has been receiving special attention in the last years, which contributed to a huge increase in its applications [47]. PCL is a semi-crystalline aliphatic polyester, rather hydrophobic, with a slower degradation rate. The main cause of PCL polymers degradation is hydrolysis. This degradation is affected by size, crystallinity of the polymer, the pH and temperature of the environment [48, 49]. Furthermore, PCL is approved by Food and Drugs Administration (FDA) for biomedical applications, has low cost and can be used in diverse scaffold fabrication technologies [47]. The major disadvantages of synthetic polymers are that their properties can differ, even for the same composition, as a function of manufacturing, temperature, sterilisation, local environment and design geometry [43]. Furthermore, they lack osteoconductivity, bioactivity and have low stiffness that is desired for bone tissue engineering [50] [51]. Notwithstanding, a polymer-based scaffold can have these properties if the scaffold contains a sufficient amount of bioactive materials such as bioceramics [52, 53].

In this way, polymer/bioceramic composite scaffolds represent a convenient alternative for applications in bone tissue regeneration due to the possibility to tailor their various properties depending on the particular needs. Composites have shown to be more effective for the enhancement of both mechanical properties and bioactivity, in comparison to ceramic and polymers alone [48, 54, 55].

1.2.2 Nanotechnological approaches for bone tissue engineering

Until few years ago, emphasis was placed on macroporous features and mechanical properties of the scaffolds. As the field of tissue engineering evolved, more importance was been given to biological aspects and to the functionalization of the scaffolds [56-58]. The importance of the reproduction of the cellular microenvironment to the regulation of essential cellular functions, such as adhesion, proliferation, morphogenesis and differentiation, showed that the reproduction of the ECM is crucial for tissue regeneration [56, 59].

Natural bone is a complex hierarchical structure of tissue and a true nano-composite of collagen nanofibers reinforced by hydroxyapatite crystals [60]. The fibrous component of the ECM provides structural support as well as surface for cell adhesion and can also regulate cell shape and migration patterns based on composition and arrangement [61]. Besides, fibrous proteins also act as storage locations for the release of bioactive molecules and growth factors upon release by proteolytic cleavage [61]. Therefore, recent efforts are being done to the production of new functional biomaterials that reproduce these hierarchical structures, though using complex chemistry and tedious procedures [62].

New bone tissue engineering methodologies and progress in nanotechnology have triggered the use of nanostructures as scaffolds for the purpose of tissue engineering [13, 63, 64]. Techniques for creating nanoscale features, patterns and particles have emerged and represent promising strategies to reproduce the natural structure and functions of the ECM. As a result, enhance communication between cells and biomaterials and promotes the desired cell behaviour [33]. Among the various nanostructures, nanofibers are very attractive for the biomedical applications, since they present a similar fibrous structure to that of natural ECM. The nanofibers can be organized into various porous architectures and possess a high surface area to volume ratio [65, 66]. On the other hand, although some nanostructures, such as nanoparticles, may have a higher surface area, nanofibers have the advantage that they can be fabricated into more sophisticated macroscopic structures such as sutures and scaffolds [65]. Indeed, the architecture and the scaffold porosity scale have an important role in cell adhesion and in cell answer to an external stimulus (figure 5) [67].

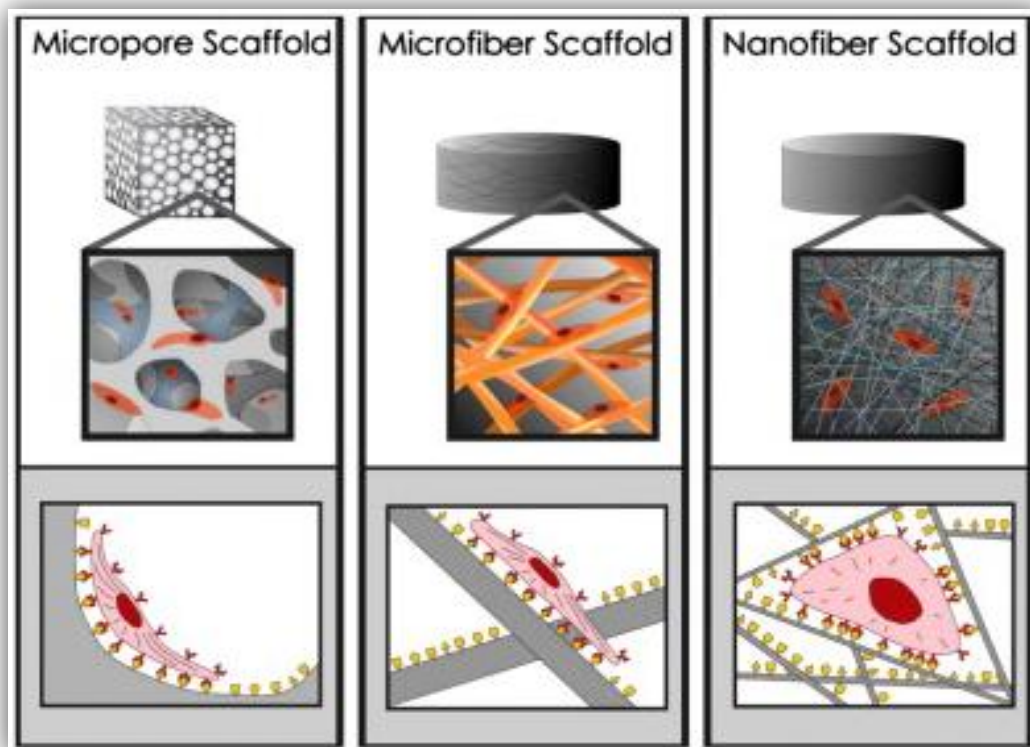


Figure 5 - Influence of architecture scale on cell binding and spreading (Adapted from [67]).

In this way, different methods for the production of polymer nanofiber scaffolds have been investigated (Table 1).

Table 1 - Comparison of different methods for the fabrication of polymer nanofiber scaffolds (adapted from [61]).

	Advantages	Disadvantages
Electrospinning	<ul style="list-style-type: none"> ✓ Easy to setup ✓ Cost effective ✓ High level of versatility, allows control over various features ✓ Vast materials selection 	<ul style="list-style-type: none"> - Poor cell infiltration into the core of the scaffolds - Toxic solvents often used
Self assembly	<ul style="list-style-type: none"> ✓ Easy incorporation of cells during fiber formation ✓ 3D pore arrangement ✓ Injectable for in vivo assembly 	<ul style="list-style-type: none"> - Complex procedure - Lack of control of fiber orientation and arrangement - Limited fiber diameter
Phase Separation	<ul style="list-style-type: none"> ✓ 3D pore arrangement 	<ul style="list-style-type: none"> - Complex procedures - Lack of control of fiber arrangement
Bacterial Cellulose	<ul style="list-style-type: none"> ✓ Low cost ✓ High yield 	<ul style="list-style-type: none"> - Limited material selection - Lack of versatility for functionalization
Templating	<ul style="list-style-type: none"> ✓ Vast materials selection ✓ Control over fiber diameter and length 	<ul style="list-style-type: none"> - Sacrificial materials - Limitation on fiber dimensions and arrangement
Drawing	<ul style="list-style-type: none"> ✓ Vast materials selection ✓ Simple procedure 	<ul style="list-style-type: none"> - Low productivity (One single fiber at a time) - Difficult to form fibers with consistent diameter
Extraction	<ul style="list-style-type: none"> ✓ Natural materials 	<ul style="list-style-type: none"> - Limited material selection - Limited control of fiber diameter and length
Vapor-Phase Polymerization	<ul style="list-style-type: none"> ✓ Polymer synthesized directly into nanofibers 	<ul style="list-style-type: none"> - Limited control of fiber diameter and length - Limited material selection - Complicated procedures
Kinetically controlled solutions synthesis	<ul style="list-style-type: none"> ✓ Polymers synthesized directly into nanofibers 	<ul style="list-style-type: none"> - Limited control of fiber diameter and length - Limited material selection - Complicated procedures
Chemical polymerization of Aniline	<ul style="list-style-type: none"> ✓ Polymers synthesized directly into nanofibers 	<ul style="list-style-type: none"> - Limited control of fiber diameter and length - Limited material selection - Complicated procedures

Nanotechnological approaches have a great potential for medical applications. In particular, the development of electrospinning is very important for health care applications since it is a relative quick, simple and cost-effective method for producing nanostructured materials desirable for many biomedical applications such as tissue engineering [65, 68, 69].

1.2.3 Electrospinning

Electrospinning has become a popular tissue engineering technique because is capable of producing fibers at the micrometer and nanometer scales, which can mimic the cellular microenvironment and therefore enhance cell adhesion and proliferation [52].

The typical electrospinning setup consists of a high power supply, syringe pump, syringe, metallic needle, and a grounded collection device [70]. In this process, a polymer solution or melt is loaded into a syringe with an attached needle [70]. The syringe pump is used to control the flow rate at which the solution is feed [71]. A high voltage is applied to the polymer solution, which overcomes the surface tension of the polymer fluid at the tip of the needle to form a charged jet [66, 72]. When the induced electrical field is above a critical value, the pendant drop at the end of the needle tip is deformed and forms a conical shape, known as the Taylor Cone [73, 74]. This phenomenon was discovered in 1969 by Sir Geoffrey Taylor and shows that a conducting fluid can exist in equilibrium in the form of a cone under the action of an electric field, where a jet can be formed at the apex of the cone, if a continuing supply of the liquid is provided [75, 76]. Due to the electrical force, the jet is accelerated. During the acceleration, however, the viscous resistance prevents the jet from moving forward, as a result, there is a decrease in acceleration [74]. In this moment, any small perturbation will prevent its straight movement, and instability occurs [74]. The three different geometric phases of the polymer fluid during the electrospinning process are illustrated in figure 6. As this jet proceeds through the air, the solvent evaporates, with the consequent dry and stretching of the polymer jet till the deposition of a polymer fiber mesh on the grounded collector [73, 77].

In the typical process, the deposited fiber mesh consists of random, non-woven and highly porous mats [59, 67]. Fiber diameter is one of the features that is very important in this structures, since the smaller the diameter of the fibers, more surface area is available for cell activities and drug loading [65]. Moreover, some authors refer that nanoscale fibers in the lack of beads, enhance performance in terms of cell response, relatively to the microscale counterpart [59]. These features, along many others, like the morphology, arrangement, pore size, surface topography and chemistry, can be controlled by varying parameters affecting electrospinning [78, 79]. The electrospinning parameters can be divided in three main groups: solution parameters, processing parameters and environment parameters [78].

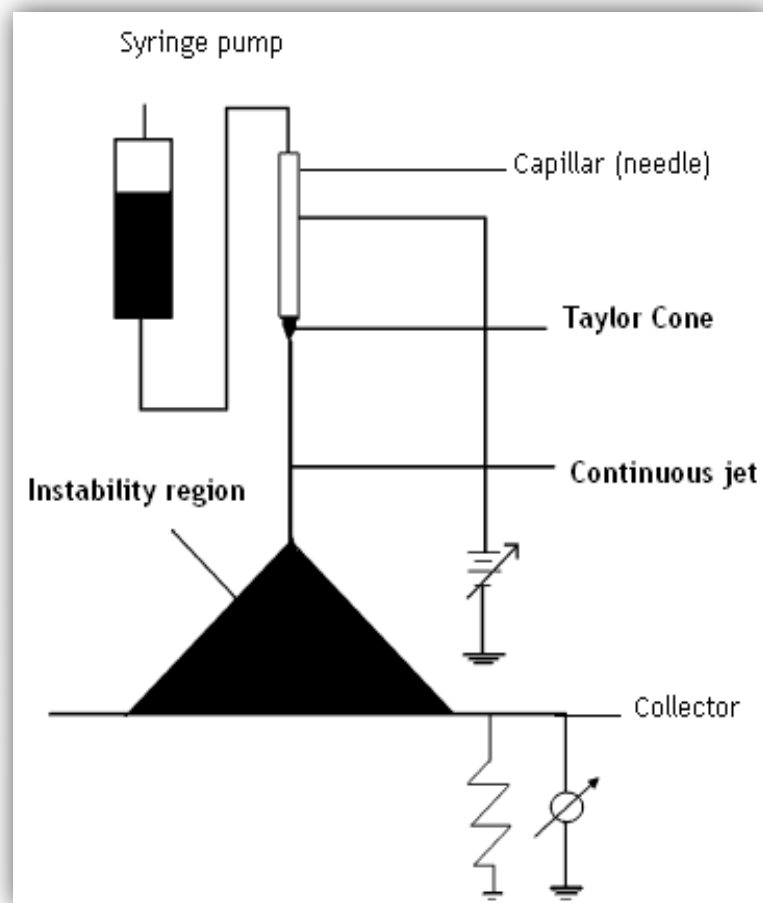


Figure 6 - Geometric phases of the polymer fluid in electrospinning process: Taylor Cone, continuous jet, instability region (Adapted from [74]).

Solution related parameters

Accordingly to what is described in the literature, tailoring the different electrospinning parameters, a multitude of natural and synthetic polymers can be used for the production of fibers at the micro and nano scales [80, 81]. The properties of the solution are the most important factors that define the electrospinnability of a polymer [78]. These properties include solution concentration, viscosity, surface tension, conductivity, dielectric constant and average molecular weight of the polymer [82, 83]. The solution concentration should be high enough to allow molecular chain entanglements [84]. If this minimum concentration is not reached, fiber formation is not achieved, instead, leads to the production of droplets when electrified, a process known as electro spraying [84, 85]. For higher concentrations the formation of beads decreases and smooth fibers can be achieved [86]. Nonetheless, the solution concentration is also directly correlated with the viscosity. Thus, a solution with too high concentration can present an excessively viscosity, preventing

the elongation of the jet and, in this way, avoids the fiber formation [86]. On the other hand, the polymer molecular weight also influences the viscosity of a solution. The molecular weight must be high enough to prevent the breakdown of the molecular chains during the electrospinning process, but not too high to not confer a huge viscosity to the solution [78, 87]. The surface tension is a resistant force that acts in the solution at the needle tip [82]. Higher surface tension increase the difficulty of the formation of a stable jet, leading to an increase of beaded structures formation [83, 88]. This can be reduced by adding a surfactant or salt to the solution [88]. This addition increases the charge density in the ejected fluid, leading to stronger elongation forces due to self-repulsion of charges under the high electrical field. In this form, thinner and more uniform electrospun fibers can be produced due to the increased conductivity [89-91]. A vital selection that influences the solution properties is the solvent used [83]. Normally, this choice is based mainly on the solubility of a polymer in the solvent. Nevertheless, this is not totally correct, given that a higher solubility is not directly related with a higher conductivity of the fluid itself [82]. A “good” solvent for electrospinning has the capability to dissolve the polymer and possesses, at the same time, a high dielectric constant. This allows the carry of a relatively bigger amount of charges, enhancing the continuous stretching of the jet, resulting in smaller diameter fibers without beads [91-93]. The dielectric constants of some of the most used solvents are presented in table 2.

Table 2 - Dielectric constants of the most used solvents applied in the preparation of electrospinning solutions (Adapted from [78]).

<i>Solvent</i>	<i>Dielectric constant</i>
2-propanol	18.3
Acetic acid	6.15
Acetone	20.7
Acetonitrile	35.92-37.06
Chloroform	4.8
Dichloromethane	8.93
Dimethylformamide	36.71
Ethyl acetate	6.0
Ethanol	24.55
m-Cresol	11.8
Methanol	32.6
Pyridine	12.3
Tetrahydrofuran	7.47
Toluene	2.438
Trifluoroethanol	27.0
Water	80.2

Process related parameters

The process parameters used in electrospinning include the electrical potential applied, the distance between the needle tip and the collector, the diameter of the needle and the feed rate (flow rate) [78, 88]. The high voltage applied to the solution is crucial for the resulting morphology of the fibers. A minimum threshold voltage is required for the generation of sufficient charges on the fluid and, subsequently, the formation of the fibers [94]. Depending upon other parameters, such as flow rate and fluid viscosity, higher voltages may be necessary to allow the stabilization of the jet [95]. In general, when the voltage increases, higher elongation forces and, thus, more stretching is imposed to the jet, resulting in a decrease of the fibers diameter produced [94, 96]. However this is not consensus, some authors refer that there is not a significantly influence of the electric field on the fiber diameter, and for higher voltages there is also a greater probability to occur bead formation [94, 96, 97]. Moreover, other authors have reported that when higher voltages are applied, more polymer is ejected which results on larger fibers diameter [94, 98]. The tip to collector distance is another parameter that tailors the fiber diameter and morphology. It is considered that this distance has to be sufficient to allow the fibers to dry before reaching the collector and to prevent the formation of a beaded morphology [94, 97]. Although it is considered not so significant as other parameters, an increase in this distance decreases the fiber diameter and for greater distances the formation of beads can also occur [94, 97]. Polymer flow rate influences jet velocity and the material transfer rate. This is very important since the flow of solution through the needle must be sufficient to replace the solution ejected and, in this form, jet be maintained [94, 97]. High flow rates may contribute to the increase of fiber diameter and an excessive flow rate augment the beads defects due to the insufficient time for the solvent to evaporate, before reaching the collector [94, 97, 99]. Other parameter that can influence the morphology of the fibers is the needle diameter. Accordingly to some authors, the fiber diameter seems to decrease by decreasing the needle diameter [96]. Moreover, Wang and coauthors have studied the effect of the needle diameter in the efficiency of electrospinning process and verified that a bigger needle diameter results in a high number of fibers produced [100].

Ambient related parameters

In electrospinning environment conditions such as temperature, humidity and air velocity in the electrospinning chamber should also be taken into account [101]. As expected, there is an inverse relationship between viscosity and surface tension of the solution and temperature [94, 101]. Moreover, solution conductivity gradually increases with temperature, thus contributing to improve the solutions electrospinnability [101, 102]. The high rate of solvent evaporation at higher temperatures also facilitates the removal of the solvent, which

may contribute to the variation of the fiber diameter [102]. The variation in humidity also has an effect in solvent evaporation rate. When humidity is very low, a volatile solvent may dry so fast that electrospinning process only is possible for little time before the needle tip is clogged [94, 102]. On the other hand, at higher relative humidity, there is a favourable adsorption of water on the polymer solution jet due to the higher partial pressure of water in the atmosphere. This does not allow the complete drying of the jet and results in fibers with pores at the surface [78, 94, 102]. The air velocity in the electrospinning chamber is also important due to the relative low mass of the fibers produced. Its spinning motion can be brutally affected by even the smallest air flow between the tip and collector [75]. Based on this fact, some authors have investigated the implementation of systems whereas the deposition and morphology of the fibers can be controlled by adjusting the air flow in the electrospinning chamber [103].

Advanced electrospinning modifications

In addition to the diverse parameters herein explained, the typical electrospinning apparatus can also be target of various modifications in order to tailor the fiber deposition and orientatio to enhance the function of the resulting fibrous structures. These modifications consist mainly on changing nozzle system (capillary) and deposition collectors [70, 104, 105]. Depending on the application, multiple and coaxial nozzles can be implemented (Figure 7).

An array of adjacent nozzles can improve fibers production efficiency at the same time that possibilities the introduction of different materials into the fibrous electrospun structure [106]. In this case, the electrical potential applied must be rigorously controlled for all the adjacent capillaries, which require special attention on the electrical field interferences between nozzles, toward the production of uniform fibers [106, 107].

In coaxial electrospinning, two polymers are ejected simultaneously from two concentrically nozzles, which results in an inner core-shell structure [104, 108]. This method is useful for the production of dual composition fibers and micro/nanotubes that can also be used for controlled drug release and bioactive tissue scaffolds [108, 109].

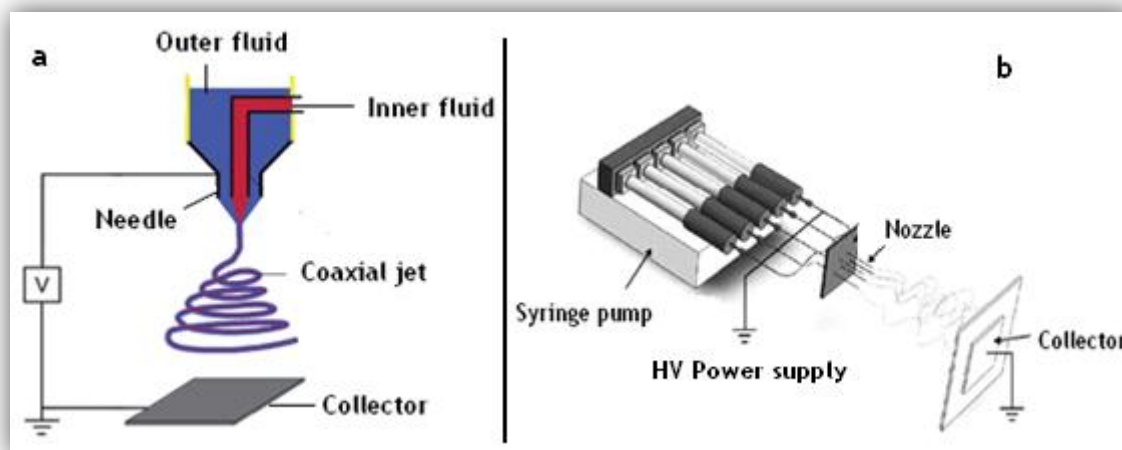


Figure 7 - Schematic setup for various nozzle types: a) coaxial electrospinning (Adapted from [110]), b) multiple nozzle electrospinning (Adapted from [106]) .

The instability of the jet results a random orientation of the fibers deposited on the collector. For many applications, the control of spatial orientation may be required. Alignment of electrospun fibers can be achieved, for example, by using a rotating drum or a pair of split electrodes as a collector, or through the introduction of an external electric field (figure 8) [105, 111].

Several studies revealed that well-aligned fibers are generated when a cylinder or disk rotating at high speed is used as collector [111]. The rotation movement increases mechanical properties through better fiber arrangement in the direction of rotation [104]. Borhani and coworkers demonstrated that when the rotation speed of the drum increases, there is an increase in the density of the nanofibers mats, resulting in a decrease of the bulk porosity of the nanofiber structures [112]. However, a higher rotation speed may also cause fiber discontinuity [70]. This method has been used for neural tissue and blood vessels engineering, where the alignment properties are essential for guiding cells migration [113-115].

The geometrical configuration of a collector has also been reported as a mean to generate uniaxially nanofibers arrays [106]. By collecting the nanofibers across the void gap of two parallel conductive collectors, highly aligned nanofibers are produced [111]. This method is very convenient when there is the need to transfer the aligned fibers onto other substrates for further processing and applications [106]. One of the major drawbacks of this method is its low productivity. This results from the fact that the most nanofibers are randomly deposited on the electrode surface instead of filling the gap [105].

In more complex setups, auxiliary electrodes can be used to create more ordered electrospun fiber structures [104]. These auxiliary electrodes may influence the deposition location of the electrospun fibers, aligning fibers and forming patterns [105]. Some authors reported that in magnetic-field-assisted electrospinning, the jet direction and diameter can

be determined by the magnetic field gradient [116]. Moreover, other authors demonstrated that by properly implementing an external magnetic field, they can prevent instabilities in the multiple nozzle electrospinning [117].

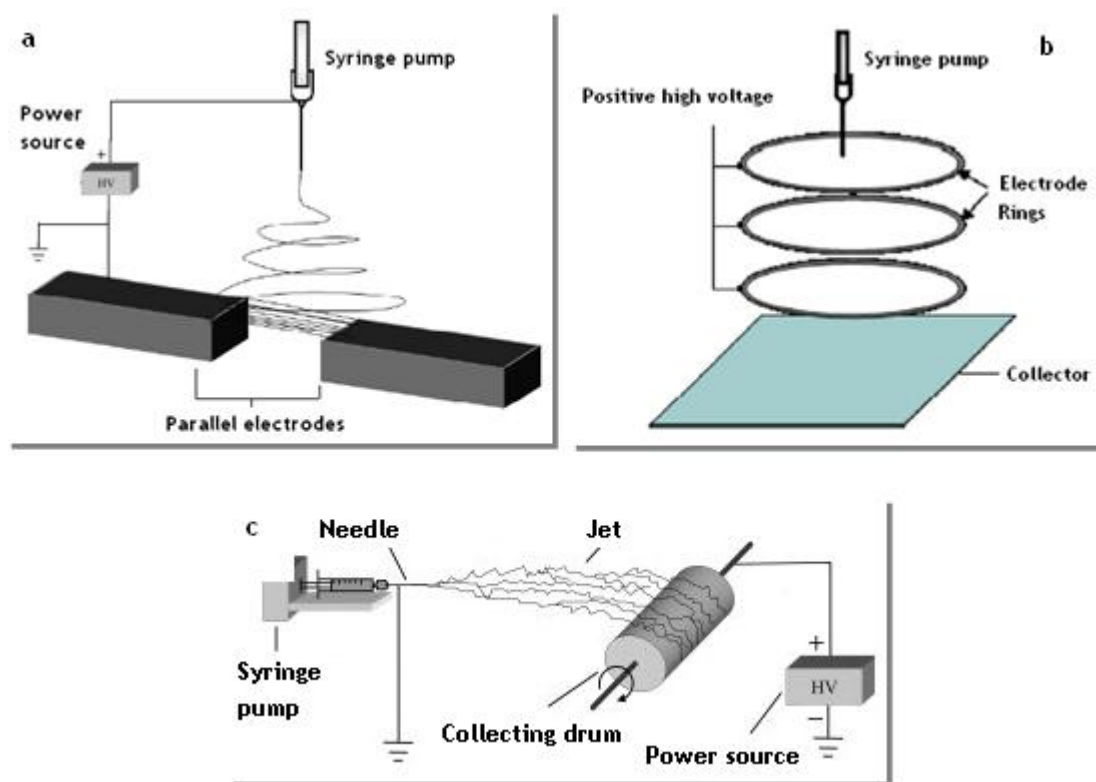


Figure 8 - Schematic illustration of various electrospinning setups for control of the fibers alignment: a) Pair of split electrodes as a collector (Adapted from [105]); b) Magnetic-field-assisted electrospinning (Adapted from [118]); c) Cylindrical rotation drum as a collector (Adapted from [119]).

1.2.4 Incorporation of growth factors into nanofibers

In order to accelerate and stimulate bone regeneration, addition of active biological molecules in the bone defect site is considered an effective and safer therapeutic method [120]. Among them, special interest has been given to BMPs, which are members of the TGF- β superfamily [121]. Of these molecules, the isoforms BMP-2, BMP-6, BMP-7 and BMP-9 are considered the most bioactive molecules for osteogenesis [120, 122, 123]. They promote migration of mesenchymal stem cells and their differentiation into osteoblasts [124].

Apart from osteoinduction angiogenesis is another essential process that is fundamental for bone regeneration and it is involved in the initiation of fracture healing. This process of neovascularisation is essential to supply nutrients, oxygen and remove products from the metabolism [125]. VEGF stimulates proliferation and migration of endothelial cells that mediate the sprout of the vascular network. Thus, the addition of VEGF in ischemic

tissue region may have positive results in bone regeneration [126]. However, when VEGF is released too rapidly or in an uncontrolled manner, various adverse effects can occur [127].

Other studies refer that BMPs and VEGF play an essential role in cell communication during osteogenesis and angiogenesis. Moreover, they report that the peak of VEGF expression occurs in the early days of bone healing and the peak of BMPs occurs at a later phase [124]. The presence and time of appearance of these signal molecules are of special relevance, since they can be used in tissue engineering approaches to improve tissue formation in vitro and in vivo [3]. Therefore, the temporal control of the tissue regeneration process is very important and it involves various agents at different times [128]. Nonetheless, a controlled delivery system that mimics endogenous growth factor production remains a challenge with conventional scaffold strategies [128].

In this work, the possibility of growth factors incorporation into nanofibers was studied. This is very important since growth factors act as cell signals that promote their proliferation and contribute for bone tissue regeneration.

1.3 The application of nanofibers for biotechnological purposes

The filtration membranes are fundamental for separation of molecules from mixtures. The membranes potential for this purpose was recognized for the first time 160 years ago, when Pfeffer and Graham were studying osmosis, and verified different rubber bands permeabilities to gases. However, the use of membranes for separation processes can be considered relatively recent [129]. Membranes can be defined as a selective barrier (selectively transfers mass between two phases) that allows a partial separation of components from a mixture, which are separated without changing their characteristics. The majority of the applications use synthetic membranes, which can be prepared either with inorganic materials like ceramics, or with organic materials including all types of polymers. There are several methods used for the production of membranes that are dependent on the membrane material and its application [129].

The separation processes using membranes can be classified based on the driving force they use to perform the separation. One of the most used processes is the one in which the driving force is the difference of pressure across the membrane. Within this, can be distinguished four types, microfiltration (MF), ultrafiltration (UF), nanofiltration (NF) and reverse osmosis (RO). The size of the solutes that membranes have the capacity to retain, decreases as it passes from MF to the RO.

In biotechnology the most common membrane technologies include operations of MF and UF. The pore size of the MF membranes varies between 0.05 μm and 10 μm , which make these membranes suitable for retention of suspended solids, emulsions and bacteria. The UF

membranes are also seen as porous; however, the pore size is considerably smaller, ranging between 10 nm to 50 nm. This feature makes these membranes mostly used to retain macromolecules. As a consequence, these membranes can be used to concentrate and purify solutions of large molecules, which comprise the major biopolymers such as DNA, RNA or proteins. The scientific research into the production of plasmid DNA (pDNA) from fermentation extracts has grown considerably in recent years, driven by the possible use of this macromolecule in gene therapy [130]. Several studies have been done on the possible use of membrane filtration in the process of pDNA purification. In particular, it is important to underline that ultrafiltration can be very useful for the purification of *Escherichia Coli* lysates, which demonstrated the feasibility of this process [131, 132].

The modification of membranes is the process by which the properties of a membrane are altered in favour of a better selectivity and / or productivity. The modification of MF membranes was done in order to prove the versatility of the electrospun PCL nanofibers. The objective was to obtain a modified membrane with a pore radius considerably smaller, in order to retain macromolecules such as pDNA.

1.4 Objectives

The combination of properties such as structural support, cellular support and controlled delivery of specific biological agents in a unique system, seems to be one way to achieve the desired efficiency in bone tissue regeneration [133, 134]. Knowing the nanotechnology potential to reproduce the extracellular environment, it may be a valuable tool for enhancing cell-biomaterial communication and promote better cellular response [33].

In the present study we used an electrospinning technique to reproduce the cellular microenvironment of the bone ECM by means of the fabrication of a PCL electrospun nanofibrous structure. Additionally, different applications for the nanofibers produced were studied. The present master thesis work plan had the following aims:

- Setup of an electrospinning apparatus;
- Optimization of the electrospinning process;
- Electrospun of PCL nanofibers;
- Incorporation of a model protein into PCL nanofibers;
- Coating 3D scaffolds with these nanofibers;
- Evaluation and characterization of the biological properties of the systems produced;
- Modification of MF membranes with the electrospun PCL nanofibers.

Chapter II

Materials and Methods

2 Materials and Methods

2.1 Materials

A high power voltage supply (Spellman CZE1000R, 0-30 kV) with very low-current output (0-300 μ A) was purchased from Spellman High Voltage Electronic Corporation (West Sussex, United Kingdom). A syringe pump (KdScientific, KDS-100 series), polycaprolactone ($\bar{M}_w=14,000$ g/mol), acetone, phosphate-buffered saline, bovine serum albumin (BSA), dulbecco's modified eagle's medium (DMEM-F12), ethylenediaminetetraacetic acid (EDTA), L-glutamine, penicillin G, streptomycin, Amphotericin B and trypsin were purchased from Sigma-Aldrich (Sintra, Portugal). 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium reagent, inner salt (MTS) and electron coupling reagent (phenazine methosulfate; PMS) were purchased from Promega. Fetal bovine serum (FBS) was purchased from Biochrom AG (Berlin, Germany). Human osteoblast cells (CRL-11372) were purchased from American Type Culture Collection (VA, USA). Pierce[®] BCA protein assay reagent A and B were purchased from Thermo Scientific. Tris Base was purchased from Fisher Scientific. Microfiltration membranes (FSM 0.45PP) were purchased from Alfa Laval (Algés, Portugal). β -Tricalcium phosphate (TCP) powder was purchased from Panreac (Barcelona, Spain).

2.2 Methods

2.2.1 Electrospinning setup

The system herein used to carry out the electrospinning process is composed of a high power voltage supply (Spellman CZE1000R, 0-30 kV), a syringe pump (KDS-100), a syringe fitted with a stainless steel blunt end needle and an aluminium plate as the conductive collector (10cmx12cm). The needle is positively charged by the power supply and the metal collector is grounded. The charged tip and grounded collector form a static electric field between them to provide the driving force that enables fiber formation [80, 94].

2.2.2 Preparation of PCL polymer solutions

PCL was dissolved in acetone under vigorous magnetic stirring, at different concentrations (7-27% (w/v)). To facilitate PCL dissolution, the various solutions were heated at 50 °C for a while and were sonicated for 1 hour [135].

2.2.3 Optimization of the electrospinning process

The previously prepared solutions were transferred into a 10 ml syringe, fitted with a blunt end needle. The flow rate of the PCL solution was controlled by the syringe pump. In this study, different electrospinning parameters were tested: electric voltage applied to generate the electrical field (10-27 kV); distance between the needle tip and grounded collector (10-20 cm); diameter of the needle (0.5-0.9 mm); flow rate (0.5-7.0 mL/h). The collection of samples for the optimization process lasted 1 min. Electrospinning was performed at room temperature [75, 135].

2.2.4 Scanning electron microscopy

The morphology of electrospun fibers was analyzed by scanning electron microscopy (SEM). Samples were air-dried overnight and then mounted on an aluminium board using a double-side adhesive tape and covered with gold using an Emitech K550 (London, England) sputter coater. The samples were then analyzed using a Hitachi S-2700 (Tokyo, Japan) scanning electron microscope operated at an accelerating voltage of 20 kV and at various amplifications [136, 137]. The diameter of the electrospun fibers was measured in a software from the microscope. During the measuring process, 2 fibers had been selected and assayed in every SEM micrograph.

2.2.5 Coating of the 3D β -Tricalcium phosphate (TCP) scaffolds with PCL nanofibers

The coating of the TCP scaffolds produced in our group with PCL nanofibers was performed by the typical electrospinning process. TCP scaffolds were placed between the needle tip and aluminium collector at a distance of 10 cm of the needle tip. A PCL solution in acetone (27%(w/v)) was transferred to a 10 ml syringe fitted with a 0.9 mm diameter needle and the electrospinning process was carried at 18 kV with a flow rate maintained by the syringe pump. For a complete covering of the TCP scaffolds, they were subject of electrospinning for 1 min, turned around, and the same procedure was repeated at 4 different positions. All the experiments were carried out at room temperature.

2.2.6 Coating of the TCP scaffolds with BSA incorporated into PCL nanofibers

PCL was dissolved in acetone at a concentration of 25% (w/v). BSA was dissolved in a small amount of acetone at a concentration of 5% (w/v). The two solutions were mixed under vigorous magnetic stirring and the resulting solution was sonicated for 1 h. PCL+BSA solution was then transferred to a 10 mL syringe fitted with a 0.9 mm needle and coating of the TCP scaffolds was performed by typical electrospinning process. Briefly, TCP scaffolds were placed between the needle tip and aluminium collector at a distance of 10 cm of the needle tip. The electrospinning process was carried with a flow rate maintained by the syringe pump. For a complete covering of the TCP scaffolds, they were subject of electrospinning for 1 min, turned around, and the same procedure was repeated at 4 different positions. All the experiments were carried at room temperature. To verify that nanofibers really had protein incorporated, TCP scaffolds coated with PCL+BSA nanofibers were transferred to a tris buffer solution. TCP scaffolds coated with PCL nanofibers were transferred at the same time to a tris buffer solution (pH=7.4) and used as controls. After one day, protein content in buffer solution was determined via BCA method on a UV-VIS Spectrophotometer (UV-1700 PharmaSpec, Shimadzu) at 570 nm.

2.2.7 Proliferation of human osteoblast cells in the presence of the different scaffolds

Human osteoblast cells were seeded in T-flasks of 25 cm³ with 6 ml of DMEM-F12 supplemented with heat-inactivated FBS (10% v/v) and 1% antibiotic/antimycotic solution. After the cells reached confluence, they were subcultivated by a 3-5 min incubation in 0.18% trypsin (1:250) and 5mM EDTA. Then cells were centrifuged, resuspended in culture medium and then seeded in T-flasks of 75 cm³. Hereafter, cells were kept in culture at 37°C in a 5% CO₂ humidified atmosphere inside an incubator [138, 139].

To evaluate cell behaviour in the presence of the scaffolds herein produced, human osteoblast cells were seeded with materials in 96-well plates at a density of 5x10⁴ cells per well, for 72 h. Previously to cell seeding, the plates and the materials were UV irradiated for 30 min [139]. Cell growth was monitored using an Olympus CX41 inverted light microscope (Tokyo, Japan) equipped with an Olympus SP-500 UZ digital camera.

2.2.8 Characterization of the cytotoxicity profile of the different scaffolds

To evaluate the cytotoxicity of the scaffolds, human osteoblast cells were seeded, at a density of 5×10^4 cells per well, in a 96-well plate, with 100 μ l of DMEM-F12 and were incubated at 37°C, in a 5% CO₂ humidified atmosphere. The plates with materials were UV irradiated for 30 min, before cell seeding. After an incubation of 24, 48 and 72 h, the mitochondrial redox activity of the viable cells was assessed through the reduction of the MTS into a water-soluble formazan product. Briefly, the medium of each well was removed and replaced with a mixture of 100 μ L of fresh culture medium and 20 μ L of MTS/PMS reagent solution. Then, cells were incubated for 4 h at 37°C, under a 5% CO₂ humidified atmosphere. The absorbance was measured at 492 nm using a microplate reader (Sanofi, Diagnostics Pauster). Wells containing cells in the culture medium without the scaffolds were used as negative controls (K-). EtOH (96%) was added to wells that contained cells, as a positive control (K+) [139, 140].

The Statistical analysis of cell viability results was performed using one-way analysis of variance (ANOVA) with the Dunnet's post hoc test. A value of $p < 0.05$ was considered statistically significant [139]. Results of cell viability in the presence of different scaffolds and in negative controls were compared with positive controls.

2.2.9 Coating of microfiltration membranes (FSM 0,45PP) with PCL nanofibers

The coating of the filtration membranes with PCL nanofibers was performed by the typical electrospinning process. The membranes were fixed on the aluminium collector at a distance of 10 cm of the needle tip. A PCL solution was transferred to a 10 ml syringe fitted with a 0.9 mm diameter needle and the electrospinning process was carried at a flow rate maintained by the syringe pump. Four different deposition times were tested: 2.5, 5.0, 7.5 and 10 min. All the experiments were carried out at room temperature.

Chapter III

Results and discussion

3 Results and Discussion

In this work an electrospinning apparatus was mounted and different assay conditions were tested in order to optimize the nanofibers production. Moreover, the PCL nanofibers were used to coat the 3D TCP scaffolds, in order to increase its surface and improve its properties desired for bone applications. Furthermore, the nanofibers were also used for decrease the pore size of MF membranes.

3.1 Electrospinning setup

The main components of a typical electrospinning apparatus were purchased and mounted as observed in figure 9. A syringe fitted with a blunt end needle was placed in a syringe pump that controlled the flow rate. The needle tip was connected to the high voltage power supply through the high voltage cable provided. The aluminium collector was linked to the ground. This is placed above a support that allows the movement for adjustment of the distance between the needle tip and the collector. To facilitate the removal of the samples, the collector plate was covered with an aluminium foil. A PCL solution was transferred to the syringe and the electrospinning equipment was tested. Initially, the typical electrospinning process was not observed. Instabilities in the electric field were responsible for this fact, since the polymeric solution was not constantly deposited on the collector surface as expected. Instead, solution splashes were thrown in all directions and not for the collector. Conducting materials like metal junctions presented in the room may have contributed to this situation. In addition, air flow may also affect the direction of the polymeric fluid under the electrical field. In this way, a plastic box was introduced in the electrospinning apparatus in order to decrease the external interferences. After this, it was noticed an improvement on the electrospinning setup and the majority of the disturbances were eradicated. However, solution splashes continued to be thrown back in direction of the syringe pump. Therefore, the incorporation of a paperboard between the needle tip and collector was carried in order to prevent this situation. The paperboard was cut with a size similar to the plastic box size and a small hole was done in the middle for needle passage. After, all the polymeric solution could be driven only for the aluminium collector, a regular deposition area appeared, as can be observed in figure 10-a, and the electrospinning setup was considered complete. It is important to underline that due to the floor electrical conductivity, a wooden pallet had to be applied in order to ensure the safety of the user, when manipulating the equipment (figure 10-b).

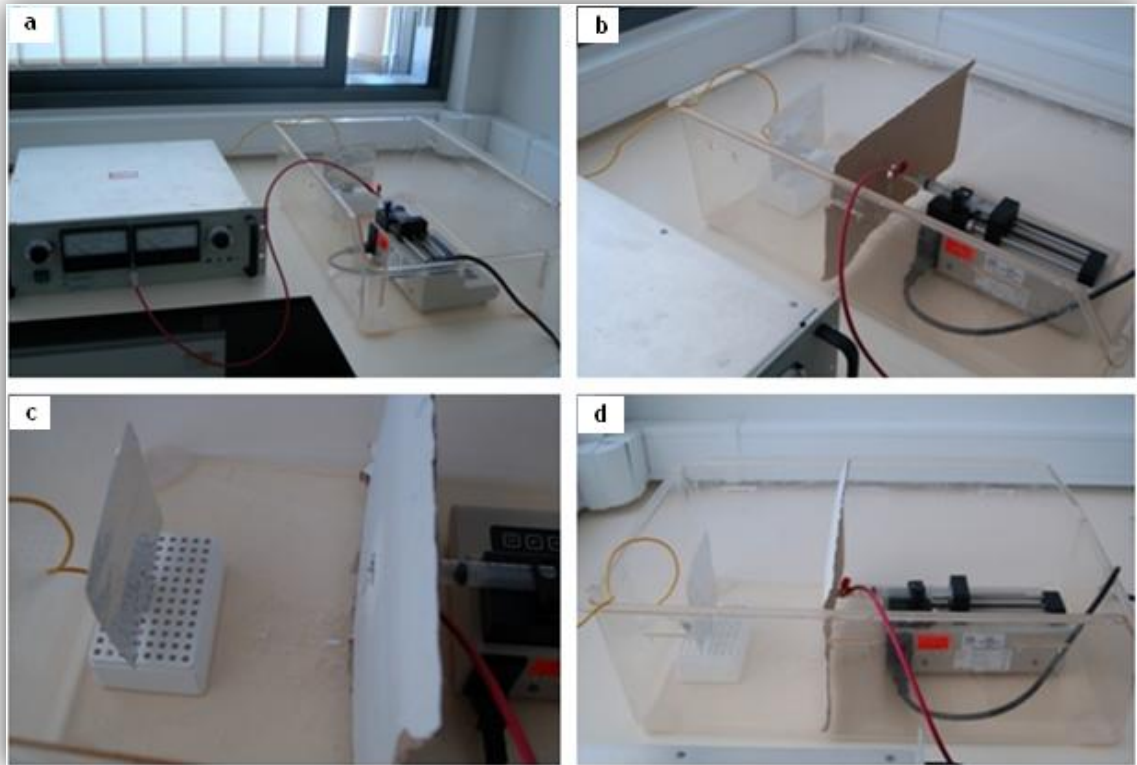


Figure 9 - Images of the electrospinning apparatus: Geral image of the apparatus (a); Paperboard with a hole for needle passage (b); Needle tip to collector distance (c); Plastic box introduced to prevent external interferences (d).

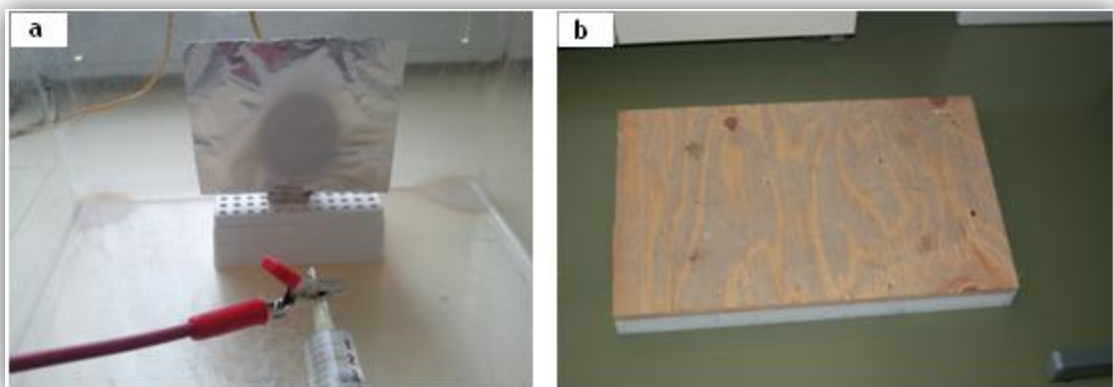


Figure 10 - Images of the typical polymeric deposition area on collector (a) and the wood pallet used to assure user's safety (b).

3.2 Optimization and characterization of the electrospun PCL nanofibers produced

As previously described in the text, in electrospinning there are a number of conditions and parameters that can greatly affect fiber production and structure. After the electrospinning setup, different solution and process parameters were tested and adapted accordingly to what is described in the literature [141, 142]. Initially, although using the parameters described in the literature, no fibers were obtained as can be seen in figure 11-a. The first SEM images revealed the formation of a polymeric cluster with irregular structure. The absence of fibers and the visualization of beads were thought to be derived from a poor solution homogeneity, or due to an incomplete chain entanglement [143]. To avoid this problem the polymeric solutions were subject to heating and sonication before electrospun, in order to facilitate PCL dissolution and enhance their homogeneity. Moreover, being acetone a good solvent for PCL and with a relative high dielectric constant [78], increasing polymer concentration was thought to solve this problem, by enhancing the chain entanglement. As explained by Shenoy and coworkers, an insufficient polymer concentration leads to formation of droplets instead of fibers, a process known as electro spraying [143]. As the polymer concentration increases and comes close to the entanglement concentration, a mixture of beads and fibers are observed. This fact was confirmed in this work, as demonstrated in figure 11-b. By increasing even more the solution concentration, there are sufficient chain entanglements to fully stabilize the jet, resulting in the reduction of jet breakup and consequently better fiber formation (figure 11-c).

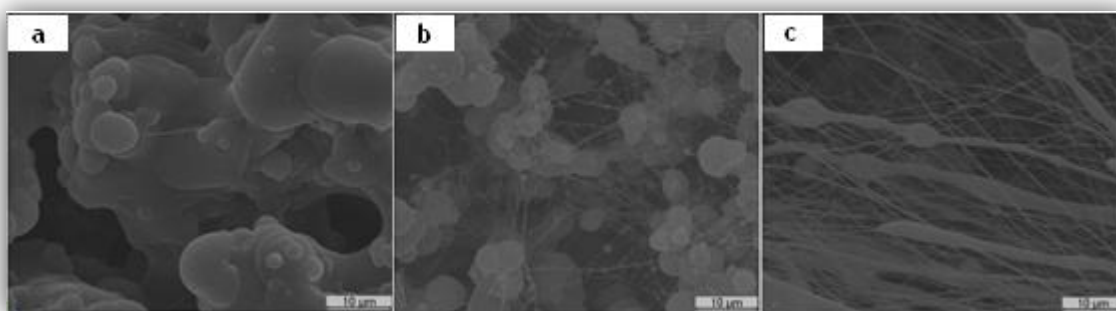


Figure 11 - SEM images of the samples obtained during the electrospinning optimization process: a) Polymeric cluster formed at low solution concentrations; b) Mixture of beads and fibers formed when polymeric concentration is approximately the same as the entanglement concentration; c) Formation of fibers for a solution concentration higher than the entanglement concentration.

Analysing the first fibers produced, it was easy to conclude that their size is situated in the nanometer range and that their morphology is still far from the desired uniformity. Therefore, knowing that several electrospinning parameters could be changed, were varied some of them to study their influence on fibers morphology. Moreover, a standardization of these results was also looked for.

In this work five electrospinning parameters were exhaustively studied: Solution concentration, applied voltage, polymer flow rate, distance between the needle and collector and needle diameter.

3.2.1 Effect of polymer concentration on nanofiber properties

The polymer concentration determines the electrospinnability of a solution, namely whether fibers are formed or not. The polymer concentration influences both the viscosity and the surface tension of the solution. Both of these properties are very important for the electrospinning process.

In this work, PCL solutions (7%, 15%, 20%, 25% and 27% (w/v)) were used to investigate the effect of polymer concentration on the nanofiber properties. All the other parameters were maintained constant to ensure that any difference observed was only caused by the variation of polymer concentration. The parameters that were kept constant were:

- . Voltage applied - 18 kV;
- . Flow rate - 1mL/h;
- . Tip to collector distance - 10 cm;
- . Needle diameter - 0.9 mm.

SEM images of the resulting samples can be visualized in figure 12. For the lower concentrations (7-20% (w/v)) the Taylor cone was not visible and a stable jet was not obtained. Consequently, there was only appearance of beads and no fibers were formed (figure 12-a). As explained before, these results may be related to the insufficient polymeric chain entanglements. Thus, too dilute solutions lead to fiber break up before reaching the collector, resulting in droplets deposition. When polymeric concentration is increased to 25% (w/v), it comes close to the entanglement concentration and a mixture of fibers and beads appears (figure 12-b). Ultimately, an increase in polymer concentration to 27% (w/v) leads to nanofibers formation (figure 12-c). These nanofibers have a slightly diameter deviation, with a mean diameter of 90 nm, approximately. Morphologically, there are also some divergences, whereas the thicker nanofibers are smooth and the narrower nanofibers tend to curl. The relative higher concentration necessary for fiber formation in this work when compared to other works, may be explained by the molecular weight of PCL. The PCL molecular weight herein used was 14,000 g/mol, whereas the typical PCL molecular weight used in electrospinning is around 80,000 g/mol [144, 145]. Then, the low polymer molecular weight herein used requires a higher concentration to achieve a sufficient number of chain entanglements for jet stabilization [78].

Beyond that, even higher polymeric concentrations were experimented to electrospun, but the needle clogged and electrospinnability was not accomplished. This fact

is also reported by other authors, whereas solutions with too high polymeric concentrations are extremely viscous, which may difficult the flow through the capillary [97].

Furthermore, in order to demonstrate if a solution electrospinnability is only dependent of the polymer concentration, lower concentrations were used to be electrospun with the variation of other parameters. It was concluded that maintaining all the other parameters constant and increasing the voltage applied to 22 kV, the Taylor cone was formed and a stable jet was observed for a 10% solution concentration. SEM images of the obtained samples revealed that nanofibers were formed and that the mean fiber diameter increased, as shown in figure 12-d.

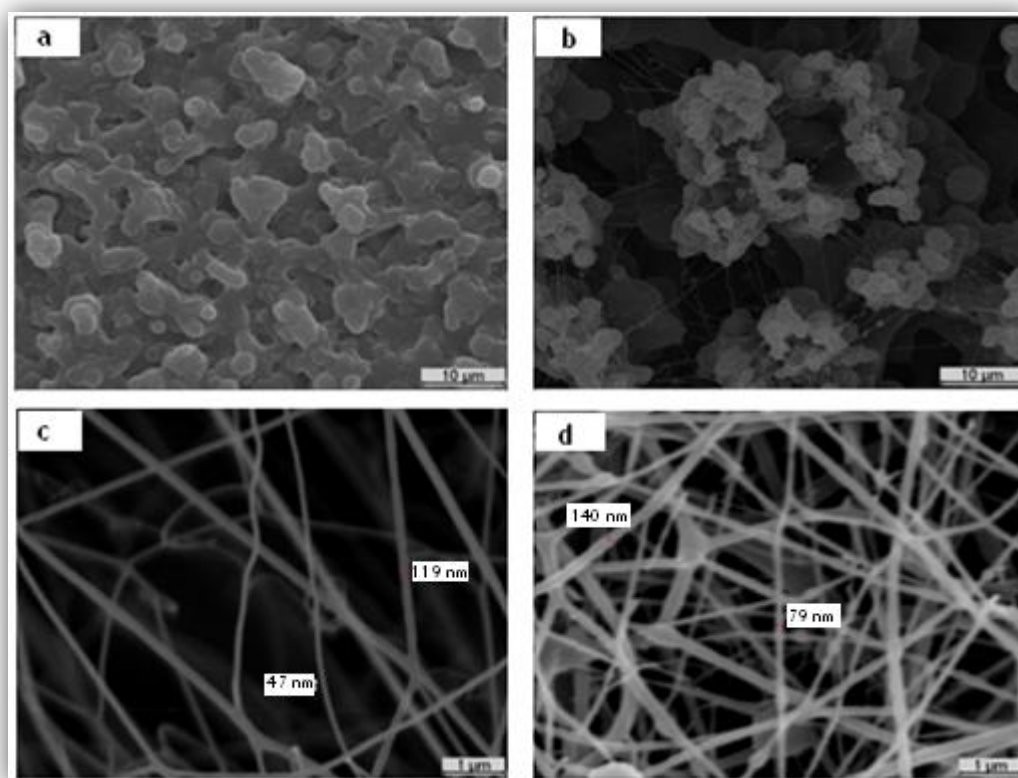


Figure 12 - Effect of the variation of polymer concentration on fibers formation: a) Image of electrospun sample with 7-20% (w/v); b) Image of electrospun sample with 25% (w/v); c) Image of electrospun sample with 27% (w/v); d) Image of electrospun sample with 10% (w/v) with augmented voltage to 22 kV and the other parameters maintained constant.

These results demonstrated that an optimum range of polymer concentrations exists, in which fibers can be electrospun, when all the other parameters are held constant. Nonetheless, polymer concentration is not the only factor that controls the solution electrospinnability. Solution electrospinnability can also be achieved by changing the voltage applied.

3.2.2 Effect of the applied voltage on nanofiber properties

In this assay, the voltage applied was changed between 10 and 27 kV, while the other parameters were kept constant:

- . Solution concentration - 27% (w/v);
- . Flow rate - 1mL/h;
- . Tip to collector distance - 10 cm;
- . Needle diameter - 0.9 mm.

According to the principle of electrospinning, a strong electric field is required to overcome the surface tension on a liquid droplet and allow jet ejection from the Taylor cone. In this work, when voltages below 10 kV were applied no Taylor cone was observed and a stable jet was not formed. Consequently, there was no fiber deposition on the collector. When voltages higher than 10 kV were applied the Taylor cone was formed and the polymeric jet seems to stabilize. SEM analysis showed that for lower voltages (10-15 kV) nanofibers with beads and with irregular shape are formed (figure 13-a,b,c). When the voltage was increased, the number of bead defects decreased until almost fully disappeared at 18 kV (figure 13-d). By applying this voltage, the nanofibers produced were relative smoother and presented the most high uniformity level of all. Furthermore, when the voltage applied was above 18 kV bead defects were observed. Moreover, the nanofibers tend to coil and some break up around the 27 kV, leaving loose ends that can be visualized in figure 13-f. Experimentally, it was observed that for higher voltages, multiple jets are formed at the needle tip. Another consequence of voltage increasing is that Taylor cone gets harder to be seen and at a critical voltage seems that it disappears. These facts were also reported by other researchers [97, 146]. Based on their works, the “disappearance” of the Taylor cone may be explained by the decrease of the volume of the pendant drop at the tip of the needle, with the increasing of the voltage. When the voltage is high enough, the jet is directly formed within the needle and no Taylor cone is seen. This is associated with higher tendency to form bead defects [97].

Additionally, it was observed that an increase in voltage did not significantly affect fiber diameter. All nanofibers presented a non-uniform diameter of around 100 nm. These results were not expected, but other authors have also reported the same results [146].

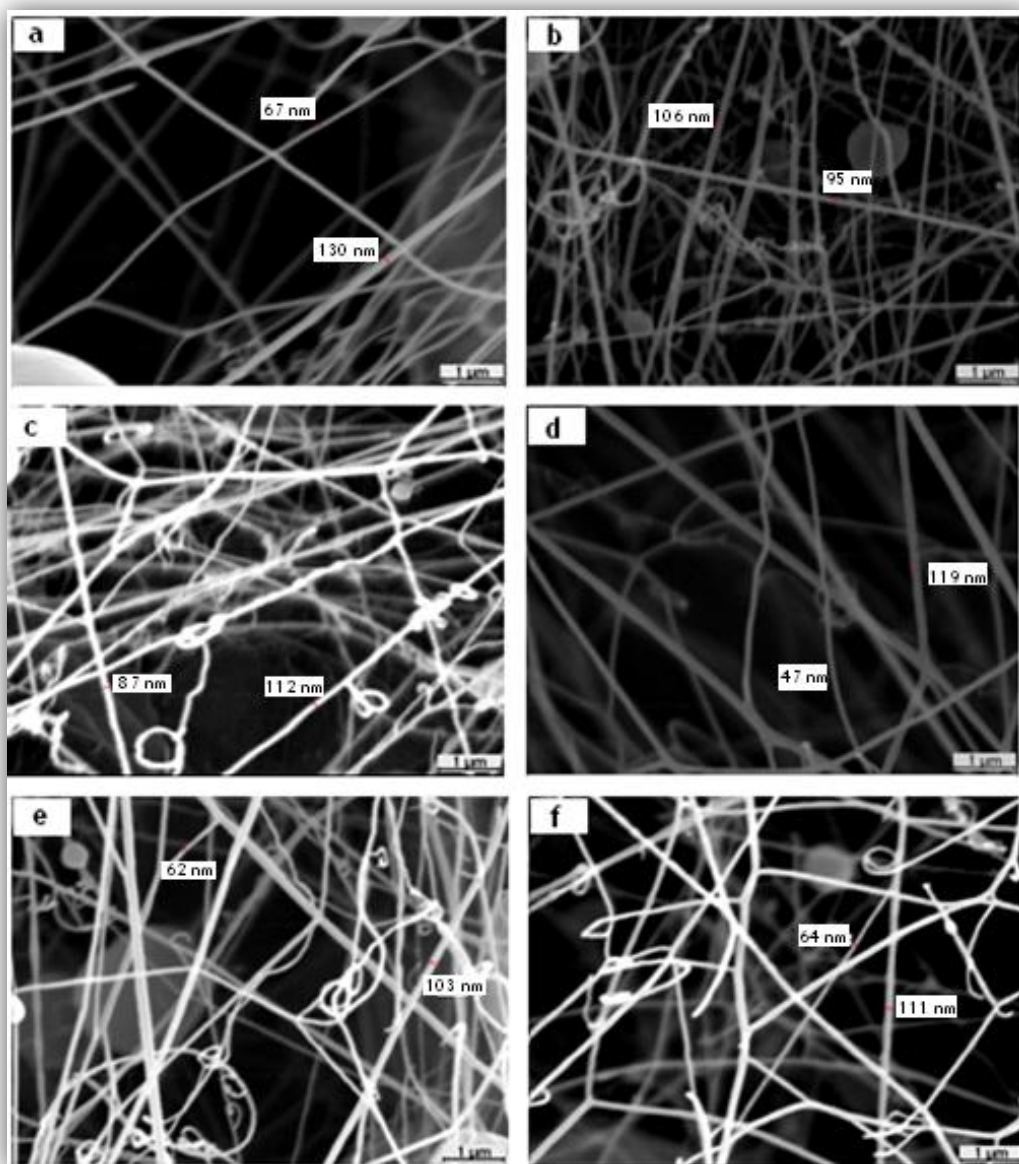


Figure 13 - Morphology of the nanofibers produced with the different applied voltage when the other parameters were held constant: a) Solution electrospun at 10 kV; b) Solution electrospun at 12 kV c) Solution electrospun at 15 kV d) Solution electrospun at 18 kV e) Solution electrospun at 22 kV f) Solution electrospun at 27 kV.

3.2.3 Effect of the polymer flow rate on nanofibers properties

In this study, the flow rate of the PCL solution was controlled by a syringe pump. The flow rate was varied between 0.5 and 7 mL/h, while the other parameters were kept constant:

- . Solution concentration - 27% (w/v);
- . Voltage applied - 18 kV;
- . Tip to collector distance - 10 cm;
- . Needle diameter - 0.9 mm.

The flow rate is responsible for maintaining a continuous and steady jet. In this work, for a flow rate of 0.5 mL/h, some nanofibers break and presented some loose ends (figure 14-a). This condition reveals that the flow of solution through the needle is insufficient to replace the solution ejected as the fiber jet [147]. A flow rate of 1mL/h seems to be the most adequate for the production of nanofibers without bead defects and with an acceptable degree of uniformity (figure 14-b). As the flow rate is increased, the formation of beads augments as well as the non-uniformity of the fibers (figure 14). This fact is explained by an excess of the delivery rate of the polymeric solution to the needle. This delivery rate exceeds the rate of solution removal from the needle tip by the electric forces and thus an unstable jet is formed along with big beads [148].

The results obtained demonstrated that nanofibers tend to have higher diameters when higher flow rates are used. Despite the divergence of the nanofibers, it is possible to see that the mean diameter of nanofibers is below the 100 nm for flow rates until 4 mL/h (figure 14-d), whereas for higher flow rates the mean diameter increases to above the 100 nm (figure 14-e).

These results suggest that flow rate has an important role on the electrospinning process and influences the morphology and diameter of the nanofibers produced.

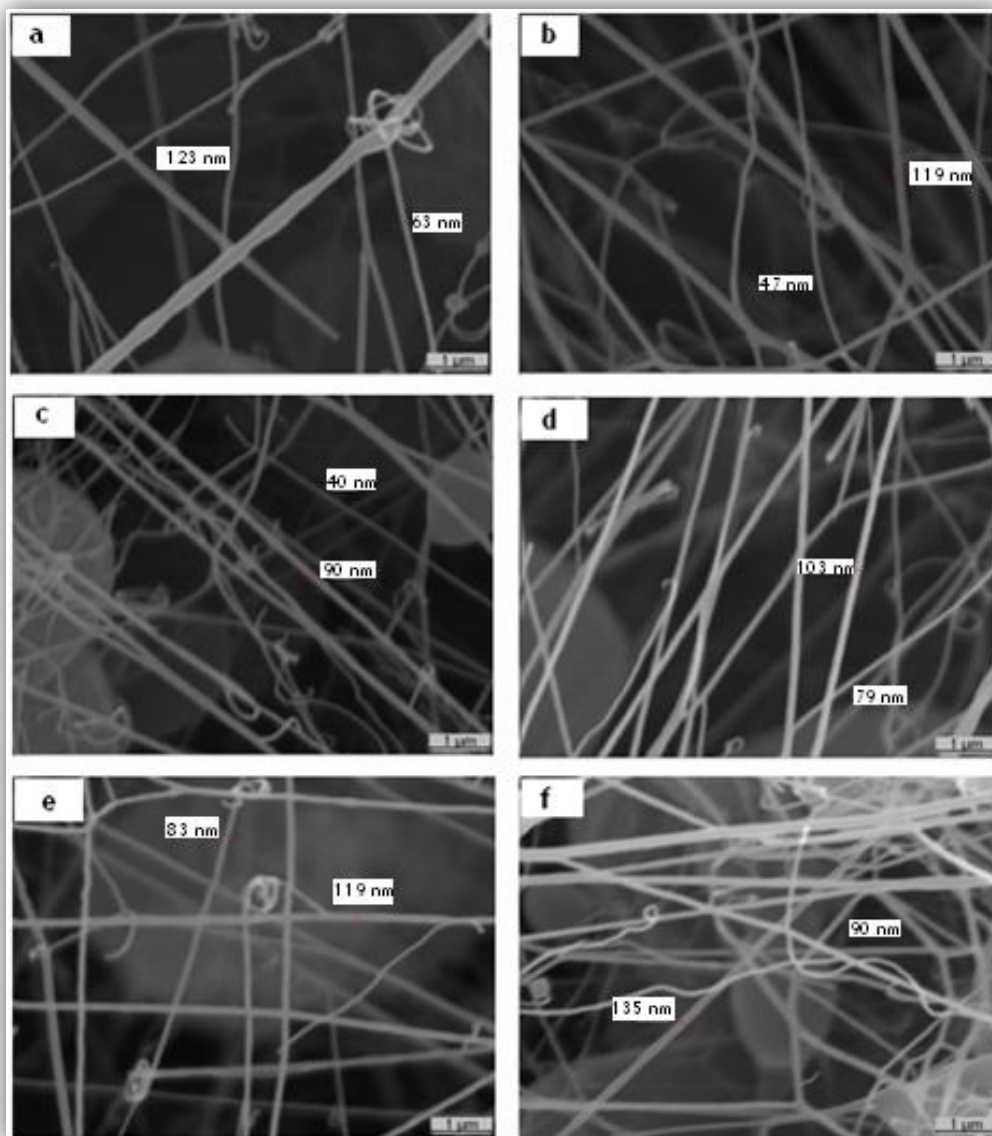


Figure 14 - Effect of the flow rate on the morphology of the nanofibers: a) Solution electrospun at a flow rate of 0.5 mL/h; b) Solution electrospun at a flow rate of 1mL/h; c) Solution electrospun at a flow rate of 2mL/h; d) Solution electrospun at a flow rate of 4 mL/h; e) Solution electrospun at a flow rate of 5 mL/h; f) Solution electrospun at a flow rate of 7 mL/h.

3.2.4 Effect of the distance between the needle tip and collector on the nanofibers properties

In this study, the needle tip to collector distance was varied between 10 and 20 cm, while the other parameters were kept constant:

- . Solution concentration - 27% (w/v);
- . Voltage applied - 18 kV;
- . Flow rate - 1mL/h;
- . Needle diameter - 0.9 mm.

The distance between the needle tip and collector has been examined as another approach to control the fiber diameter and morphology. The analysis of SEM images from figure 15, showed that all the resultant nanofibers have a similar morphology when varying the distance between the tip and the collector. This fact is in agreement with the literature, whereas some authors refer that the tip to collector distance do not have so much importance as other parameters to the fibers morphology [149].

However, it was observed a decrease in the nanofiber diameter when the tip to collector distance was increased. This happens because for larger distances, solvent has more time to evaporate, resulting in a bigger stretching of the fibers [149]. However, a tip to collector distance of 20 cm seems to have no influence in the fiber diameter and nanofibers become thicker (figure15-d). This situation may occur due to the tip to collector distance be too large, which is responsible for weakening the electric field and thus influences the electrospinning [150].

On the other hand, a slightly increase in bead defects is observed for greater tip to collector distances and some nanofibers break up are noticed (figure 15-c). This may result from an excess of solvent evaporation due to a longer time for this process to occur, which may cause too many jet splits [149].

The results obtained show that the optimum distance between the tip and collector favours the evaporation of solvent from the nanofibers and, in this case, it seems that this distance should be below the 15 cm.

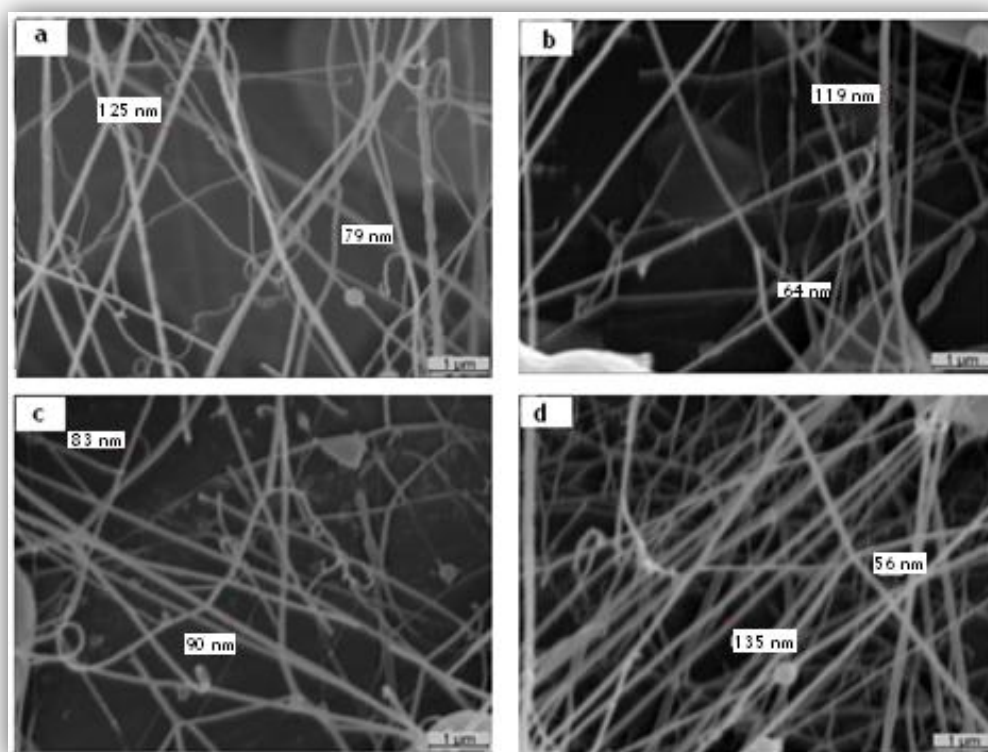


Figure 15 - Effect of the distance between the needle tip and collector on the morphology of the nanofibers, when all the other parameters are held constant: a) Solution electrospun at a needle tip to collector distance of 10 cm; b) Solution electrospun at a needle tip to collector distance of 13 cm; c) Solution electrospun at a needle tip to collector distance of 15 cm; d) Solution electrospun at a needle tip to collector distance of 10 cm.

3.2.5 Effect of the needle diameter on the nanofibers properties

The needle diameter was also changed between 0.5 mm and 0.9 mm, while the other parameters were kept constant:

- . Solution concentration - 27% (w/v);
- . Voltage applied - 18 kV;
- . Flow rate - 1 mL/h;
- . Tip to collector distance - 10 cm.

The influence of the needle diameter on the morphology of electrospun PCL nanofibers was examined experimentally. A needle with an inner diameter of 0.5 mm was tried to be used for electrospun a PCL solution. Unfortunately, this was not possible since the needle clogged and, consequently, no polymeric jet was formed. The needle opening seems to be too small and do not allow the polymer solution to flow. This may be due to the high

acetone volatility, what dries the pendant drop on the needle surface and subsequently blocks the solution passage. Based on this, larger needle diameters should facilitate the flow through the needle opening. This was confirmed since a diameter of 0.6 mm already allowed the formation of nanofibers (figure 16-a). However, small instabilities still occurred since for this needle diameter nanofibers with bead defects were formed. On the other hand, for larger needle diameters (0.8 and 0.9 mm) no problems occurred and smooth nanofibers were formed, in particular with the 0.8 mm diameter needle (figure 16-b).

In this work, the variation of the needle diameter did not affect the diameter size of the nanofibers. This is in agreement with other authors results that reported that there is no correlation between the needle diameter and the average nanofiber diameter obtained in the solution electrospinning process [151].

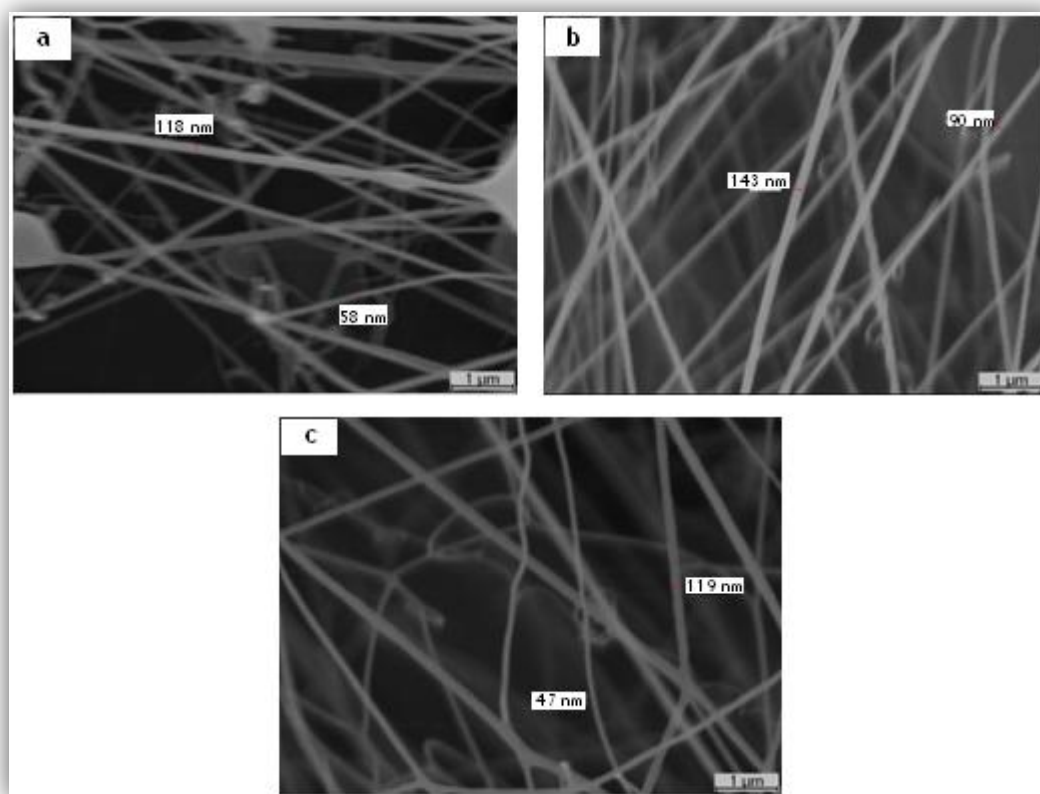


Figure 16 - Effect of the needle diameter on the morphology of the nanofibers, maintaining the other parameters constant: a) Solution electrospun through a needle diameter of 0.6 mm; b) Solution electrospun through a needle diameter of 0.8 mm; c) Solution electrospun through a needle diameter of 0.9 mm.

3.3 Coating of the TCP scaffolds with PCL nanofibers

As previously described in the text, tissue engineering is highly depended on the properties of the scaffolds. A crucial point for a scaffold to be successful applied in bone tissue engineering is the combination of mechanical properties and biological activity, since all of them play a critical role in cell response and tissue regeneration. Calcium phosphate materials have been widely used as bone substitutes in orthopedic and reconstructive surgery, due to their biocompatibility, osteoconductivity and good bone integration [39, 152].

The PCL nanofibers herein produced present high porosity, pore size and fiber dimensions that are suitable for adhesion, spreading and migration of cells. Thus, combining the better physical properties of ceramics with the good biological properties of the PCL nanofibers appears a good strategy for bone tissue regeneration. In this work, we also performed the coating of 3D TCP scaffolds with PCL nanofibers, in order to enhance biological performance of these scaffolds. Furthermore, the incorporation of a model protein (BSA) was also investigated as a preliminary assay to evaluate the system capability for controlled drug delivery. TCP scaffolds were successfully coated; nanofibers adhered well to the surface of the TCP scaffolds and a highly porous mesh of smooth fibers was formed (figure 17 - a). Moreover, BSA did not withdraw the electrospinnability of the PCL solution. Instead, PCL+BSA nanofibers looked even more uniform than the solely PCL nanofibers (figure 17 - b). In addition, when the BCA method was used it was determined a higher absorbance for the TCP scaffolds coated with PCL+BSA nanofibers than for the control (TCP scaffolds coated with PCL nanofibers). This preliminary assay suggested that this system may have potential to be used as a controlled drug delivery system.

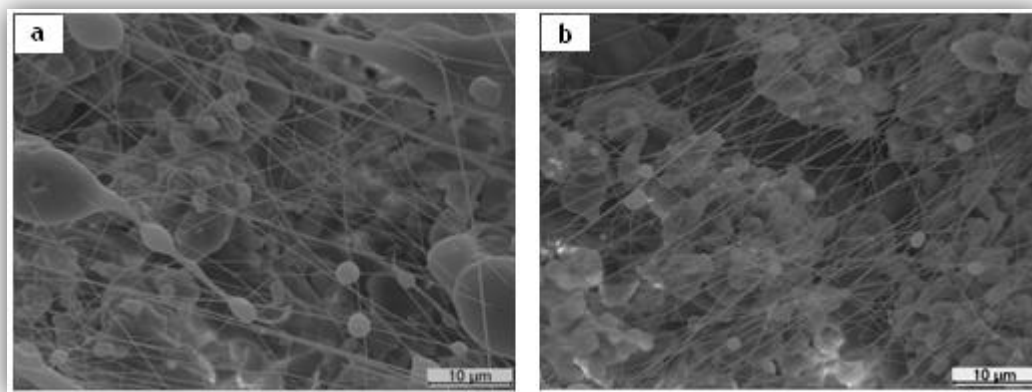


Figure 17 - SEM images of the coated TCP scaffolds: a) TCP scaffold coated with PCL nanofibers; b) TCP scaffold coated with PCL+BSA nanofibers.

3.4 Evaluation of the cytotoxic profile of the different materials

The cytocompatibility of PCL nanofibers, TCP scaffolds and TCP scaffolds coated with PCL nanofibers was characterized through *in vitro* studies. As previously mentioned, human osteoblast cells were seeded at the same initial density in the 96 well plates, with and without materials to assess its cytotoxicity. In the first 24 hours, cell adhesion and proliferation was noticed in wells where cells were in contact with the materials (figure 18, A-G) and in the negative control (figure 18 - J). In the positive control, no cell adhesion or proliferation was observed. Dead cells with their typical spherical shape are shown in figure 18 - N.

After 48 hours, cells continued to proliferate in wells where cells were in contact with the materials and in the negative control. However, it was observed a relative decrease on cell proliferation for TCP scaffolds (figure 18 - H), when compared with the other materials (figure 18 - B,E) and with the negative control (figure 18 - L). In positive control no proliferation was observed (figure 18 - O).

After 72 hours, cells in wells containing the TCP scaffolds presented the lowest proliferation (figure 18 - I). Cells in wells containing the PCL nanofibers presented high proliferation (figure 18 - C) closer to the negative control (figure 18 - M), followed by cells in wells containing TCP scaffolds coated with PCL (figure 19 - F). As expected, no proliferation was observed in the positive control (figure 18 - P)

The observation of cell growth in the presence of materials during 72 hours demonstrated that cells in contact with PCL nanofibers presented the higher proliferation, similar to that observed in the negative control. On the other hand, cells in contact with TCP scaffolds had the lowest proliferation. The relative increase on cell proliferation for TCP scaffolds coated with PCL nanofibers, when compared to that observed for TCP scaffolds solely, indicates that coating materials with PCL nanofibers can bring benefits in terms of biocompatibility.

To further evaluate the cytotoxic profile of the materials, MTS assay was also performed. The MTS assay results (figure 19) showed that cell viability was higher for the negative control, in which cells were seeded just with DMEM-F12. Cells seeded in the presence of PCL nanofibers showed that cell viability was maintained over time. Cells seeded in the presence of the other materials (TCP and TCP+PCL) showed that in both cases cell viability decreased over time, mainly after 48 hours. However, this decrease was more noticeable for cells in the presence of TCP scaffolds, when compared with TCP scaffolds coated with PCL nanofibers. As should be expected the positive control showed almost no viable cells.

The MTS assay showed a significant difference between positive control and the negative control and cells exposed to materials over 72 hours of incubation ($*p<0.05$). These results demonstrated that PCL nanofibers do not affect cell viability over time. Furthermore,

coating TCP scaffolds with PCL nanofibers improve biological properties of the TCP scaffold solely.

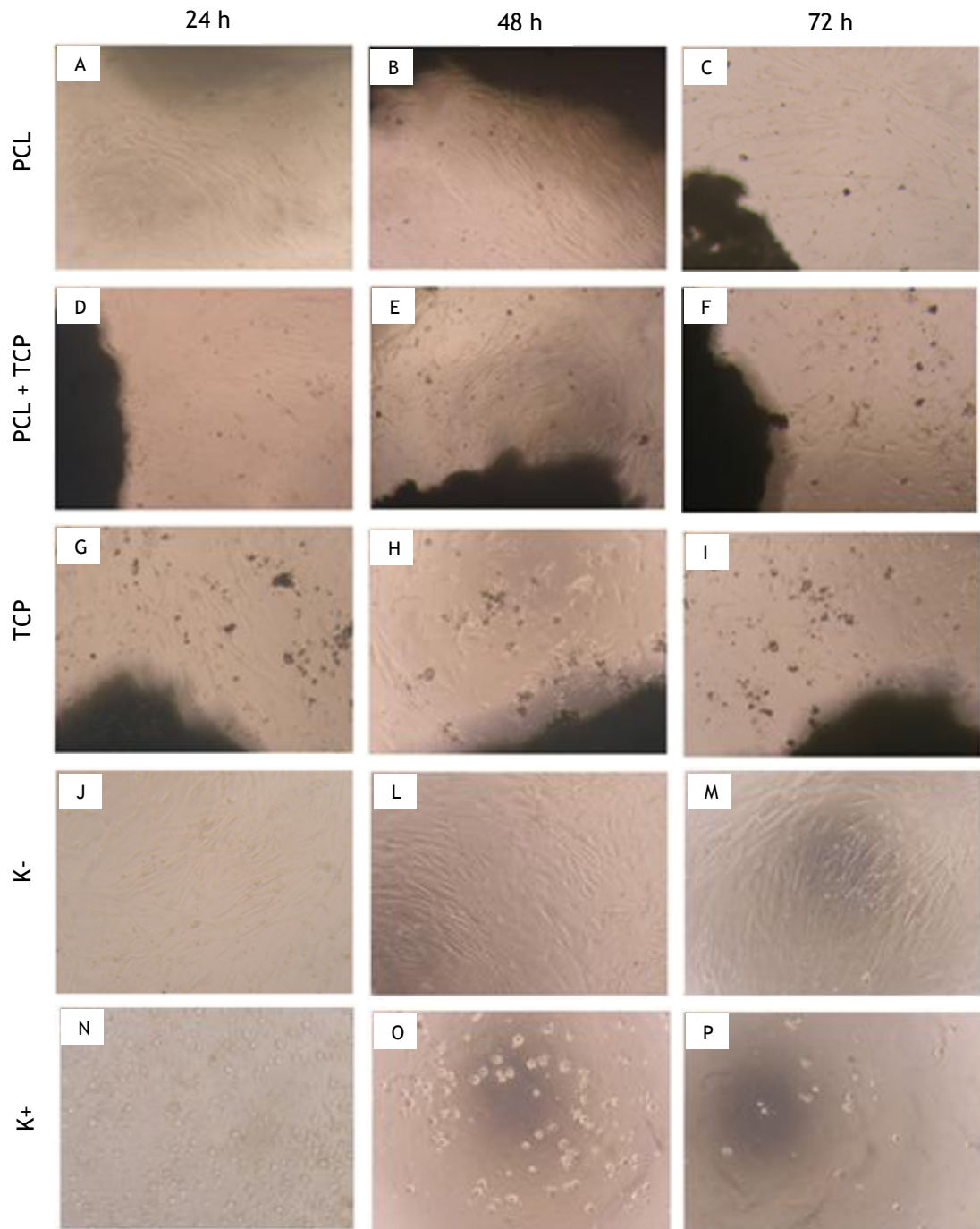


Figure 18 - Optical microscopic photographs of human osteoblast cells after 24, 48 and 72 h of being seeded. (PCL) cells seeded in the presence of PCL nanofibers; (PCL+TCP) cells seeded in the presence of TCP scaffolds coated with PCL nanofibers; (TCP) cells seeded in the presence of TCP scaffolds; (K-) negative control; (K+) positive control. Original magnification x100.

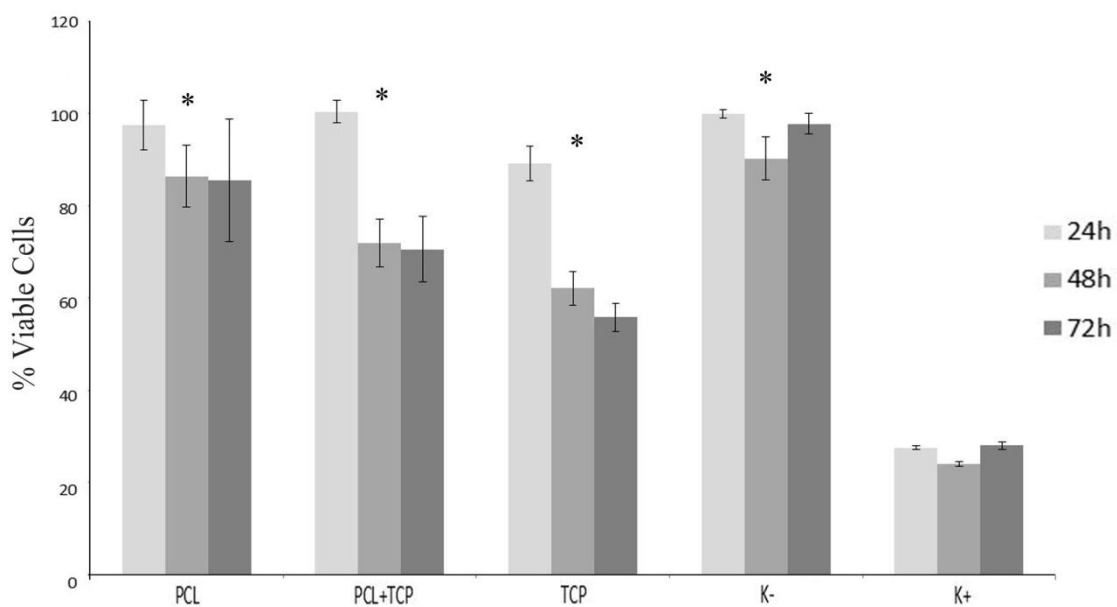


Figure 19 - Cellular activities measured by the MTS assay after 24, 48 and 72 h of being seeded. (K+) positive control; (K-) negative control; (PCL) cells seeded in the presence of PCL nanofibers; (PCL+TCP) cells seeded in the presence of TCP scaffolds coated with PCL nanofibers; (TCP) cells seeded in the presence of TCP scaffolds. Each result is the mean standard error of the mean of at least three independent experiments. Statistical analysis was performed using one-way ANOVA with Dunnet's post hoc test ($p < 0.05$).

3.5 Modification of microfiltration membranes (FSM 0.45PP) with PCL nanofibers

As stated before, filtration membranes are used in biotechnological applications to concentrate and purify diverse products resultant from fermentation processes. A pressure across the surface of the membrane creates a driven force that forces any components smaller than the pore size of the membrane to pass through. Any components larger than the pore size cannot pass through and are retained. In this work, microfiltration membranes were modified by coating them with electrospun PCL nanofibers in order to verify the effect in the retention of pDNA. Initially, the pore size of the FSM 0.45PP microfiltration membranes was 0.45 μm and when the nanofibers were added, this pore size decreased (figure 20). It was easy to see that PCL nanofibers were smaller in diameter than the membrane pores and, thus, different deposition times of nanofibers could decrease the pore size at different degrees. In this context, four deposition times (2.5, 5.0, 7.5 and 10 min) were tried to evaluate the variation in the retention of pDNA. The results of the pDNA retention for the different deposition times (n=3) are represented in figure 21. The normal retention of the commercially available membrane FSM 0.45PP is represented at deposition time 0 min. The results demonstrated a gradual increase in the retention of the pDNA until 5 min of nanofiber deposition. This increase was almost 50% for 5 min of nanofiber deposition into membrane surface, when compared with the commercially available one. For higher deposition times (7.5 and 10 min) the pDNA retention decreased and the inherent error increased. This variability in the results may be explained by a too long deposition time of nanofibers. In this way, the layer of nanofibers became too thick and could not contain itself stucked at the membrane surface when a pressure was applied for the filtration process. Consequently, membranes partial coating was lost, what resulted in lower pDNA retention values with a bigger inherent error.

Nevertheless, given the high rejection of pDNA for coated membranes during 5 min, these results are promising regarding the concentration and purification of the pDNA or other macromolecules.

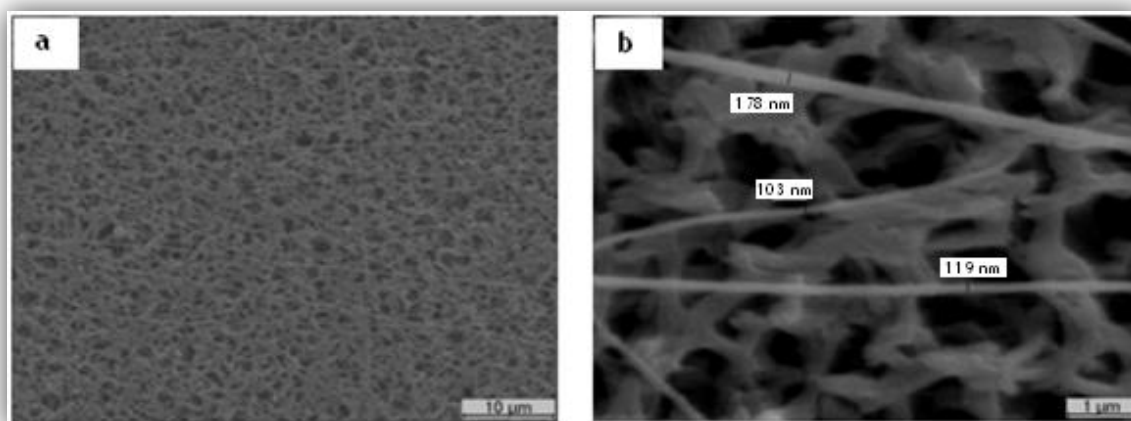


Figure 20 - Commercial microfiltration membranes coated with PCL nanofibers: a) Original magnification x2000 b) Original magnification x15000.

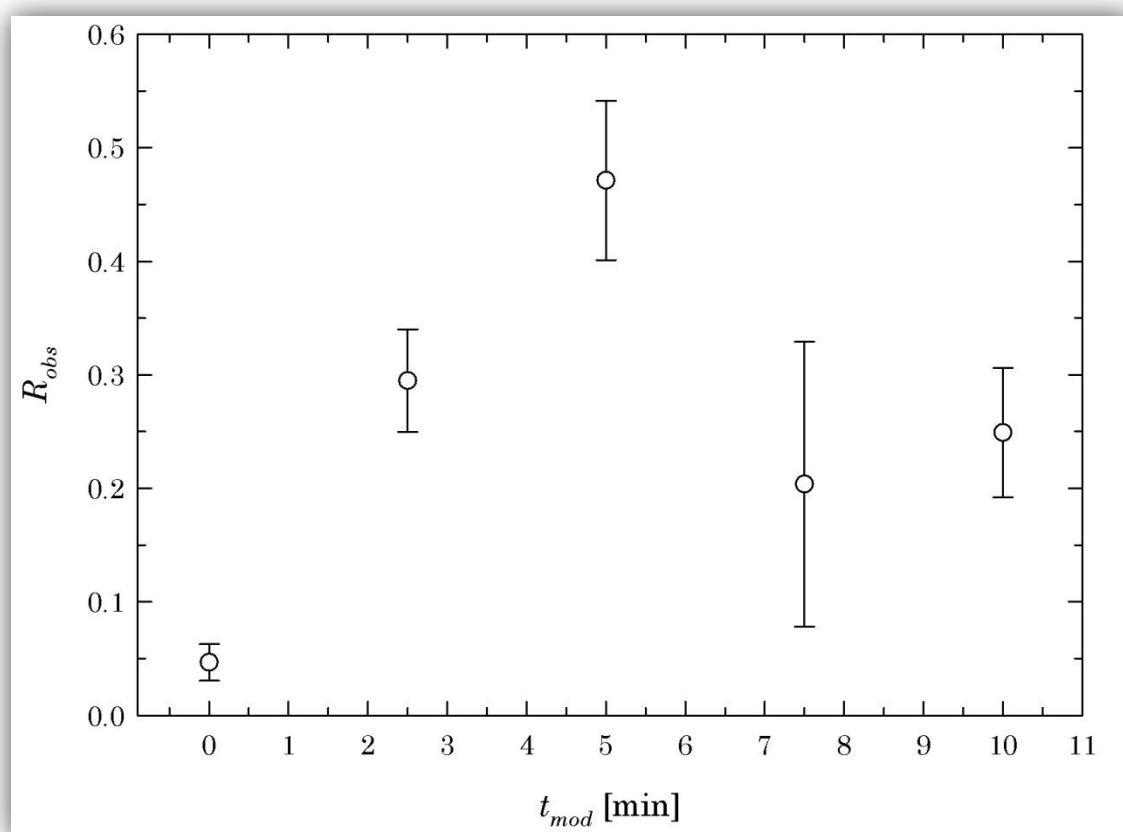


Figure 21 - Retention of pDNA for modified MF membranes (FSM 0.45PP) in function of the deposition time of the electrospun PCL nanofibers.

Chapter IV

Conclusions and future perspectives

4. Conclusions and future perspectives

Tissue engineering is a vast field that uses diverse tools in the search for successful tissue regeneration. In this context, this MsC dissertation thesis describes a method to mimic the nanostructure of bone in order to enhance bone tissue regeneration. To accomplish this, an electrospinning setup was prepared and the production of PCL nanofibers was done. During the optimization process different parameters were experimented, which influence nanofibers morphology. Although diverse morphologies were observed, nanofibers diameter was maintained between 40 nm and 170 nm. Due to the poor physical properties of this nanofibrous structures, the combination of these with ceramic scaffolds, which possess better mechanical properties, was tried. The coating of TCP scaffolds with PCL electrospun nanofibers was achieved and demonstrated an increase in the biological performance of these systems, when compared to TCP scaffolds alone. The observation of human osteoblast cells proliferation in the presence of the different materials and the MTS assay results corroborated this fact. Moreover, the incorporation of a protein model (BSA) into the PCL electrospun nanofibers was accomplished, without compromising the electrospinnability of the polymer solution. This preliminary study points to the possibility of growth factors incorporation into this system, which may accelerate bone tissue regeneration.

Based on these facts, future works may and should be developed in order to improve the system capabilities for bone tissue regeneration. The production of electrospun composite nanofibers appears as one viable option to mimic not only the fibrous nanostructure of bone, but also to mimic the hydroxiapatite crystals within it. Additionally, incorporation of nanoparticles with encapsulated growth factors into the electrospun nanofibers will allow a more protected and controlled growth factor release. These studies will, eventually, enhance biological performance of the system and consequently improve bone tissue regeneration.

Furthermore, the versatility of the herein produced PCL nanofibers was demonstrated for a different application. The modification of MF membranes was performed by coating them with the PCL electrospun nanofibers. This study revealed a huge increase in the membranes rejection of pDNA, when compared with the commercial MF membranes. Despite the good results obtained, the future coating of MF membranes with, in this case, negative charged nanofibers may enhance even more the rejection of negative charged pDNA, due to the repulsion forces. Charged nanofibers can be achieved by changing the solvent or polymer of the electrospun solution. Hence, the modified membranes demonstrate great potential for concentration of pDNA in biotechnological applications.

Electrospun PCL nanofibers, which were the base for this work, demonstrate great potential for diverse biomedical applications due to their favourable properties as high porosity, nanometer size, biocompatibility and relative ease of combination with other materials and structures.

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