# Novel hybrid membrane of chitosan/poly ( $\varepsilon$ -caprolactone) for tissue engineering

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We investigated the potential use of 3D hybrid membrane: poly ( $\varepsilon$ -caprolactone) (PCL) mesh using rotary jet spinning with subsequent chitosan (CH) coating. The morphological examinations by scanning electron microscopy (SEM) were proved the efficiency of this technique on obtaining relative homogeneous PCL fiber mats (15,49 ± 4,1µm), with high surface porosity (1,06 ± 0,41µm) and effective CH coating. The feasibility of rotary jet spinning allowed the solvent evaporation during the process; this fact was verified by differential scanning calorimetry (DSC), indeed also had verified changes in thermal properties on the hybrid membrane, since the present of CH. It was investigated the mechanical properties of the hybrid membrane and CH film, the data were that the samples presents good tensile modulus but low strain at the break. In addition, it was verified the biocompatibility properties in vitro using Vero cells. PCL mesh demonstrated cells more spread vastly in the pore surface, with attachments in between fibers indicating the potential for cell adhesion. The films samples (CH and hybrid membrane) resulted in a cells layer on the surfaces with an intense staining (metachromasy), which is the result of cells more active. The cell counting -5 days of culture- and the MTT assay -21 days of culture- demonstrated that the materials tested proved to be different from the positive control and equal to each other and this fact, in our view, this indicates a satisfactory proliferation. Thus, based on the results here, this novel hybrid membrane provides an attractive material for tissue engineering applications.

### Introduction

Chitosan (CH)-based membranes seem to be excellent materials that could be very used in many biomedical applications. This biopolymer ensures not only biodegradation, but also its natural characteristic of being an antimicrobial agent and guaranty biocompatibility with cell culture. However, less flexibility in regulating the mechanical properties limits its usage.<sup>1</sup>

Indeed the polymer poly ( $\varepsilon$ -caprolactone) (PCL), well-known semicrystalline aliphatic polyester used extensively and FDAapproved for biomedical applications, presents highly crystalline structure that is beneficial to its mechanical properties. But it has low surface energy and hydrophobicity limits its biocompatibility and has a slow degradation rate (up to 6 to 36 mo).<sup>2</sup>

The search for ideal biomaterials is still on-going for tissue regeneration where the properties of scaffolds are dictated concurrently by many factors, e.g., cell-material interaction, mechanical solicitation, degradation rate and metabolic route. Blending polymers is an approach to develop new biomaterials exhibiting combinations of properties that cannot be reaching by individual polymer. Indeed, the research of CH and PCL blend brings to a new direction for tissue engineering.<sup>3-5</sup>

In tissue engineering scaffolds can be produce by many technologies. With the focus to reach nanofibers the technique widely used is electrospinning, which can be classified by the method of polymeric preparation into solution and melt electrospinning. Since 1934, many researches attempt to improve the melt and solution electrospinning drawbacks. The solution electrospinning process suffers from low productivity (up to 300mg/hr); requirement of additional solvent extraction process; and environmental concerns (toxic solvents are used). In other hand, the melt electrospinning is free from those drawbacks, but present difficulties inherent in finer fiber formation, higher viscosity of molten polymer and the electrical discharge problem associated with the application of high voltage to polymeric melt.

The search to overcome the use of electrostatic force lead to the creation of different methods such as meltblowing, drawing, biocomponent spinning, forcespinning, phase-separation, and flash-spinning. Indeed all the methods present advantages and disadvantages comparing with electrospinning. Simple processes, free solvent use, high production rates and environmental advantages are examples of the mains vantages; however the nonuniform fiber size, higher variation of fiber diameter, complex process and machine, and polymeric heating are the principals' drawbacks.

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Figure 1. SEM images: (A-B) PCL fibers by rotary jet spinning-scale 10µm and 2µm; (C-D) Hybrid membrane- scale 20µm and 10µm.

The rotary jet spinning (RJS) was created in 2010 by Badrossamay. This system fabricates three-dimensional aligned nanofibers by exploiting a high-speed rotating nozzle to form a polymer jet which undergoes stretching before solidification. Some aspects can be controlled by varying rotational speed nozzle geometry and solution properties e.g., fiber diameter, morphology and web porosity. Comparing with the most common method to obtain fibers the electrospinning, the RJS has several advantages, e.g., no requirement of high voltage, fiber fabrication is independent of solution conductivity, can be applicable to polymeric emulsions and suspensions and high productivity.<sup>6</sup>

This study explored the application of the PCL mesh produce by rotary jet spinning technique (RJS). The aims of this research were to investigate the potential use in tissue engineering of the 3D hybrid structure: Chitosan coating/Poly ( $\varepsilon$ -caprolactone) mesh (hybrid membrane).

#### Results

### Characterization of the samples SEM images

We produced using the technique rotary jet spinning PCL fibers in a morphology of non woven membrane showing high porosity surface the average was  $1.06 \pm 0.41 \mu m$  (Fig. 1A and B).





Also, analyzing the SEM images was possible to note the full coating of the biomaterial with chitosan (Fig. 1C and D). The size of the PCL fiber were determinate by the use of the software Image Tool, a total of 54 fibers were measured at 3 randomly selected places, the average was  $15.49 \pm 4.1 \mu m$ , as demonstrated in the histogram (Fig. 2).

#### Thermal properties

PCL is a crystalline polymer: pure PCL melts at 60 °C and its glass transition temperature around -60 °C.<sup>7</sup> On the other



Figure 3. DSC Thermogram of PCL mesh and hybrid membrane PCL/CH. In the up corner is show the crystallization process and the main figure is show the second heating scan.

Table 1. Tensile modulus, peak load and strain at break of the CH and hybrid membrane samples

Samples	Tensile Modulus (MPa)	Peak load (N)	Strain at break (%)
CH film	2836,92 (± 276)	24,06 (± 1,5)	1,88 (± 0,31)
Hybrid membrane (CH/PCL)	1342,25 (± 187)	16,58 (± 3,5)	2,82 (± 0,82)

hand, the biopolymer chitosan is an amorphous biopolymer, the transition temperatures may vary significantly based on deacetylation degree reach for the used CH.

Figure 3 showed the second heating thermograms for a control PCL mesh and the hybrid membrane PCL/CH. Into figure is shown the melting peak for each sample and the enthalpy involved during the melt process. On the upper corner on the graphic is shown the crystallization process for each sample were is clear the difference between the peak crystallization temperature reach after erase the thermal history.

#### Mechanical tensile test

The samples were tested in a dry state. Table 1 shows the results the samples were very brittle, exhibiting a break strain as low as 1.8-2.8% and an elastic modulus of 2836-1342 MPa. All the samples had a uniform thickness of  $14 \pm 0.08 \mu m$  and  $12 \pm 0.03 \mu m$  hybrid membrane and CH film, respectively.

# In vitro tests

Figure 4 we observed cells growing on the materials with 1 and 2 d of culture with light microscopy. Figure 5 shows we could see cells culture by 3 d and stained with TB and CB. We see that the spacing between the material fibers interfere with the cell adhesion and growth on biomaterials. The cell growth analysis showed that after one day of culture, the positive control was different from the other samples. After two days of incubation, the sample PCL mesh was similar to negative control and both were different from others samples studied. With three days of culture, the biomaterials tested were shown to be equal to each other and different from controls.

During the culture period cells were not able to reach confluence on the substrates (Fig. 6). In all samples, at different time, we detected cells strongly stained by TB, which are basophilic dye. On PCL mesh, we found ortochromatic cells. On the other hand, on hybrid membrane and CH film we found metachromatic cells. The cells were also stained with CV, a general dye used for morphological studies. With this dye we could see cells with irregular morphology on biomaterials.

Figure 7 showed the absorbance obtained from an MTT assay of Vero cells with were cultured on chitosan films, PCL films and hybrid membrane for 21 d. The results indicate that Vero cells had viability pattern similar to positive control used. Thus, after 21 d of cultured, we found on polymers studied a number of living cells similar to culture plate.

#### Discussion

The development of new technologies to manufacture nanofibers and microfibers (e.g., drawing out, molecular self



Figure 4. Images with 24 (A1-C1) and 48 (A2-C2) hours of culture, images of the optical microscopy without staining, magnification of 100x (A-PCL mesh; B-hybrid membrane; C-CH film).

assembly, thermally induced phase separation, electrospinning) are increasing nowadays. Modifications of electrospinning techniques are co-axial electrospinning and electrohydrodynamic printing, indeed those processes use needle for the polymer ejection. Needleless technology is a promising process, since it is very flexible and enables the creation of nanofibers with high production capacity on an industrial scale, the process use liquid surfaces on a rotary spinning roller or wire. However the utilization is limited by the major disadvantages like low production rate and low safety features.

The need of improvement techniques attempted to new method without application of high voltage, in this case, centrifugal spinning system (rotary jet spinning) offers several appealing features such as the obtained fibrous web, with interconnected pores, facile and low cost effective process.<sup>8</sup> In this study we investigate the use of a Poly (caprolactone) (using



Figure 5. In vitro test Vero cells: Images with staining TB (A1-C1) and CV (A2-C2) with 72 h of culture, magnification of 200x (A- PCL mesh; B- hybrid membrane; C-CH film).

chloroform as a high volatile solvent) mesh by rotary jet spinning process with Chitosan (dissolved by acetic acid) coating for tissue engineering applications.

The solvent choice brings others possibilities at the methodology, e.g., the evaporation and viscosity control. The volatility of the solvent act at the solidification and contraction of

the jet, therefore when use highly volatile solvents the jet form thicker fibers.<sup>6</sup> Also, the fiber diameter is correlated with the dynamic viscosity, that increase proportional.<sup>9</sup> The PCL fibers had the diameters ranging from 11.39 to 19.59  $\mu$ m (Fig. 2). The literature demonstrated results lower than we obtained; with an orifice diameter of 340  $\mu$ m, 12000 rpm rotation speed and 10



**Figure 6.** In vitro test Vero cells: Cells numbers x days of culture. Black line -negative control, red line- positive control, Blue line- PCL mesh, Green line- PCL film, Pink line - hybrid membrane and light blue line- CH film. Statistically differences (P > 0.05) compared by one-way ANOVA.

wt% PLA in CHCl<sub>3</sub> had fibers with diameters of 833 to 2168 nm<sup>6</sup>; 30000 rpm rotation speed and 6 wt% PCL in1,1,1,3,3,3-Hexafluoro-2-propanol presents samples with 250 to 550 nm<sup>9</sup>; and 2000 rpm rotation speed and 12 wt% PCL in CHCl<sub>3</sub> had fibers diameters with an average of 311nm.<sup>10</sup> The research with lower rotation speed had a modification at the sample collector; the samples were collected on a round bottom collector, which provides tension to the fibers, creating densely packed, highly aligned nanofibers. In the same study was used PVP to increase the formation of surface pores which was found to be 375nm. We obtained pores in a range of 0.65 to 1.47µm (Fig. 1A and B), due the solvent evaporation. Indeed the effect of solvent on the surface structure of the fiber is an important factor is allows the increase of the surface topography, which could influence the cell adhesion and subsequent proliferation.<sup>11</sup>

PCL has a glass transition temperature of -60 °C, a melting temperature of 58-60 °C, and a decomposition of 350 °C.12 Also in the literature is reported that the PCL blended with CH improves water absorption capability, which is desirable property in many biomedical applications. Figure 3 demonstrates the main peaks of PCL mesh and for the polyester in the hybrid membrane PCL/CH. This thermograms shown the absence of the solvents used and we can compare the PCL phase in each case. During the crystallization process, when the CH is presence a heterogeneous crystallization process is inducing because the peak of crystallization temperature (Tc) is bigger in +6 °C than pure PCL. This heterogeneous process occurs during the control cooling process in the DSC apparatus but if it is occur is because some chemical affinity in the melt was produce between PCL and Chitosan. After the crystallization process, in the main figure, the second heating is shown and the difference between melt temperatures is around 3 °C, been bigger for the PCL in the hybrid sample because the heterogeneous crystallization

(that was explained before). But the nucleation process not only change the melt temperature if also change the crystallization degree, because the presence of the chitosan phase not let to PCL reach its original crystallization degree and this can be justify for the difference in the melt enthalpy show for PCL and PCL/CH.

PCL is a versatile synthetic polymer that can be mechanical altered by regulating its crystal structure. Its bulk mechanical properties include a tensile strength of 16 MPa, tensile modulus of 400MPa, flexural modulus of 500MPa, elongation at yield of 7%, and elongation at failure of 80% for PCL with Mw 44000.<sup>12</sup> Due the mechanical properties this polymer is very used as blends, composites and copolymers. <sup>3</sup>CH has long been considered as one of the most attractive natural biopolymer matrices for bone tissue engineering due to its structural similarity to the glycosaminoglycan found in bone and biocompatibility property. On the other hand less flexibility in regulating the mechanical properties and biodegradation limits are the main disadvantages of CH. Indeed the development of biomaterials with programmable mechanical and biological properties will significantly help tissue regeneration technologies.

Table 1 summarizes the mechanical properties of the samples (tensile modulus, peak load, and strain at break). The hybrid membrane composed of PCL and CH had higher strain at break which is important to ensure dimensional stability properties to this system against external forces. But the high stretching caused by the rotary jet spinning technique over the non woven PCL fibers, coupled with the presence of layers of chitosan (a very rigid but fragile biopolymer), is what gives it these low properties to this hybrid membranes.

Cytotoxicity tests were conducted to investigate the effects of the biomaterials on animal cells, which is one of the prerequisites for implantation.<sup>13</sup> The **Figure 4** showed the samples PCL mesh, hybrid membrane and CH film during 1-d and 2 d of culture, also it was possible to observe the morphology of the Vero cells, all the samples presents a compatibly surfaces, without any toxicity.

In order to see the growth and morphology of the cells on the surface were staining with toluide blue (TB) and crystal violet (CV) at 3-d of culture (Fig. 5). With CV it was seen that the cells had different interactions with different material structures. The sample with fibers demonstrated cells more spread vastly in the pore surface, with attachments in between fibers indicating the potential for cell adhesion. The films samples (CH and hybrid membrane) resulted in a cells layer on the surfaces. Cytochemistry revealed changes in cell behavior induced by materials. TB-stained basophilic cells were observed in all samples. At pH 4.0, TB stains nucleic acids and glycosaminoglycans. Although Vero cells synthesize glycosaminoglycans, they produce them in soluble form in the culture medium.<sup>13</sup> Thus, the presence of variations on basophilic cytoplasm suggests variation on cell active. We found a more intense staining (metachromasy) at cells on hybrid membrane and CH film than PCL mesh. These results suggest that cells were more active on hybrid membrane and CH film.

Once attached, Vero cells proliferated on all types of samples and were quantified by counting cell number. The 1-d results were without difference of the proliferation rate, except by negative control. With two days incubation, the sample PCL mesh was similar to negative control and both were different from others samples studied. With three days of culture, the biomaterials tested were shown to be equal to each other and different from controls (**Fig. 6**).

The culture plates, used as negative control, are prepared to stimulate cell growth. Accordingly, it was expected that these samples had higher cell proliferation. The materials tested proved to be different from the positive control and equal to each other. In our view, this indicates a satisfactory proliferation standard. Similar results were obtained with Vero cells with different blends of PLLA/PHBV and different porous PLLA scaffolds.<sup>14,15</sup>

With the goal to investigate the cell viability of the samples with Vero Cells, we produced the cell culture during 21 d, we made a MTT assay with this time of incubation. The results obtained with samples chitosan, chitosan / PCL and PCL shown that these did not alter the proliferative index, because the cells grew and multiplied on them in a pattern similar to positive control (**Fig.** 7). These data support the interpretation that the cells grow in a satisfactory manner on these polymers. The bioresorbable materials have a good ability to support cell growth. In this sense, our data are consistent with other reports.<sup>15</sup>

# 1.0 0.9 0.8 0.7 Absorbance (540nm) 0.6 0.5 0.4 0.3 0.2 \*\* 0.1 0.0 Control + Control -Chitosan Chitosan/PCL PCL Samples studied

**Figure 7.** Absorbance of the Vero Cells during 21 d of culture. The samples analyzed were Control +, Control –, Chitosan film, PCLmesh with Chitosan coating (hybrid membrane) and PCL film (\*\* P < 0,001).

Figure 8. Simulation of the rotary jet spinning equipment.

cover with the PCL mesh, after 24 h was added the 50% of the CH/acetic acid solution to form a uniform coating (Fig. 9).

#### Fabrication of the films

Like a control a film of PCL was casted into a glass petri dish by dissolving in chloroform (10% w/v) at room temperature. The dried film was collected and vacuum dried for 48 h. The same preparation method was used for CH film, but the solvent used was  $0.3_{\rm M}$  acetic acid solution with the CH/solvent proportion of 0.17:50% w/v.

#### Characterization

#### Thickness and fiber dimensions

The thicknesses of all samples were measured using a micrometer and the average value was computed from 6 measurements of different scaffolds for the same sample batch. The PCL fiber dimensions were characterized using Image

# **Materials and Methods**

#### Materials

The polymer (poly(D-glucosamine) name chitosan (CH) with viscometer molecular weight 130.000 g/mol and 80 KDa polycaprolactone (PCL) were purchased from Sigma Aldrich, the solvent chloroform from Merck and acetic acid form Synth.

#### Formation of the mesh

PCL pellets were dissolved in chloroform at a concentration of 10% (w/v) under stirring. After the homogenization, 6 h, the solution was prepared to use at the rotary jet spinning system (Fig. 8).

The equipment consisted of a reservoir with four side wall orifices, with the diameter of 0,5mm that is attached to the shaft of a motor with one rotation speed (3500 rpm). To facilitate the fiber collection an aluminum foil is placed on the cylindrical collector held against the collector wall at a distance of 300mm. The process allows obtain fibers through a drop by stretching a component of centripetal force (tangential) when from a container in rotation, is ejects a viscoelastic solution.<sup>16</sup> This is how on the inner walls of the drum, a film type non woven mesh is created.

#### Fabrication of hybrid membrane

Layer by layer CH/PCL/CH was prepared using the PCL mesh, obtained by rotary jet spinning, and solution of CH/acetic acid. The 50% of the solution was placed into a Teflon mold and



Tool software (UTHSCSA- Version 3.0), the average value was determinate using the measurements of 54 fibers of the same sample.

## Scanning electron microscopy

The surface morphology of the films were characterized using Cambridge Stereoscan 200 (operating mode, high vaccum, secondary electron SE detector) - the samples were gold coated, using the machine Sputer Coater (Bal-Tec SCD 050) with 40 mA,  $5 \times 10^7$  Pa for 200 s.

# Differential scanning calorimetry

Thermal properties of samples were characterized by Metter-Toledo (FP 85 TA Cell) differential scanning calorimetry analyzer. Around 10 mg of hybrid membrane were sealed in aluminum pans and heated up to 100 °C and maintained at the temperature for about 5 min in order to erase the thermal history of the samples. They were cooled to -40 °C from where they were heated back to 100 °C. The melting temperature of PCL was taken as the temperature at which the endothermic peak occurred in the second scan. For the CH samples it was heated up to 180 °C and maintained at the temperature for about 5 min, then were cooled to 25 °C from where they were heated back to 300 °C.

# Tensile mechanical tests

Tensile tests of samples were performed at MTS tensile tester (FEM-Unicamp). Rectangular strips (~7 mm  $\times$  ~0.5 mm  $\times$  ~0.15 mm) were subjected to a constant strain rate (20 mm/ min) in tension until broken. From the stress-strain curves, tensile modulus (MPa), peak load (N), and strain at break (%) were determined. The reported values are an average of five measurements.

# In vitro tests

Vero cells, a fibroblastic cell line established from the kidney of the African green monkey (*Cercopithecus aethiops*), were obtained from the Adolfo Lutz Institute, São Paulo, Brazil. Vero cells are recommended for studies of cytotoxicity and cellsubstratum interactions in biomaterial research.<sup>13</sup> The cells were cultured in 5% CO<sub>2</sub> at 37 °C in 199 medium (Lonza Group Ltd) supplemented with 10% fetal calf serum (FCS, from Nutricell Nutrientes Celulares).

A cell suspension  $(3 \times 10^5$  cells/well) were inoculated in medium 199 with 10% FCS at 37 °C with 5% CO2 on different biomaterials analyzed (PCL and chitosan films, PCL mesh and hybrid membrane). With 1, 3, and 5 d of culture were captured images of cell on wells with inverted optical microscope (Olympus IX71, with 4× and 10× of magnification) using *Infinity* software. At 5 d of culture, the samples were also fixed in 10% formalin (in PBS 0.1 M, pH 7.2), washed twice in distilled water, and stained with Toluidine Blue at pH 4.0 (TB) and Crystal Violet (CV). After staining, samples were mounted and analyzed under a light microscope (Nikon, model 80i), with the 40× and 10× objective lens, obtained using Nikon digital camera (DS-R1') and *NIS Elements* software. Were obtained from 4 to 12 images for each sample, were the same field was photographed at 10× objective every 0.5  $\mu$ m apart and 40×

objective, images were captured every 0.2  $\mu$ m. Captured images were merged in *Combine ZP* software and processed on *Axio Vision* software. As a negative control, we used the same cell inoculation into wells without biomaterials. As a positive control, ethanol added to the wells where we had growing cells.

The cell viability was investigated by the MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide; thiazolyl blue) assay. The samples chitosan films, PCL films, and hybrid membrane were fixed in each bottom of a 96-well cell culture plate and kept under UV light during 20 min. The different sterilized samples (n = 6), were inoculated with culture medium for 24hs at 37 °C in a 96 wells culture plate. After this incubation time,  $100\mu$ l of cell suspension ( $1.0 \times$ 10<sup>4</sup> cell/ml) in 199 medium (Lonza) with 10% FCS (Nutricell) was inoculated in the wells with different samples. Cell-free wells with the same culture conditions were used as control reactions. The cells were cultured for 21 d at 37 °C, washed twice with 0.1M phosphate buffered saline (PBS) in pH 7.4, at 37 °C, and then received 100µl of FCS-free medium and 10µl of 3-(4,5-dimetiltiazol-2-il)-2,5 difenil bromete tetrazolium (MTT, by Sigma). After 4hs in dark, 100µl of sodium dodecyl sulfate (SDS, Merck) was added. After 12hs, the wells were read in microplate reader (Microplate Reader DNM 9602, Beijing Perlong New Technology Co, Ltd.) with 570nm wavelength. As a positive control the culture plate itself (polypropilene) was used while alcohol 70% served as negative control. We also read the absorbance of all experimental conditions in a cell free condition for control of MTT reaction.

# Statistical analysis

The significant differences were analyzed by one-way analysis of variance (ANOVA). Differences were regarded as statistically significant at P < 0.05.

# Conclusions

In this paper, we present data which show the extent of the potential of the PCL fibers, produced by rotary jet spinning, coated with chitosan forming a 3D hybrid membrane with high potential to be use in tissue engineering. It was investigated the surface morphology, thermal and mechanical properties, and biocompatibility of the novel hybrid biomaterial.

The rotary jet spinning was investigated and successfully used to produce PCL 3D porous mesh. Also, the technique of casting promoted the combine of PCL mesh with CH membrane. Finally, the successful addition of cells is creating and even more versatile and biocompatible 3D hybrid structure. This preliminary investigations indicate that these strategies have great potential to improve current biomaterials and in the development of new scaffolds for tissue engineering.

#### Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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