Synthesis of poly (lactic acid) and production of scaffolds by electrospinning

Abstract

In this research, the synthesis of L-lactide (L-Lc), the cyclic dimer of poly (L-lactic acid) (PLLA) with a 55.20% yield and its polymerization to PLLA with different molecular weights is reported. The product was used to produce fibrous scaffolds by electrospinning using variations in concentration, distance, flow and voltage. A high-resolution stereoscope was employed to find fiber diameter (500 ± 1000µm); PLLA with 295,000g/mol molecular weight (viscometry) was obtained after 96 hours of reaction and yielded the best results in the electrospinning process. These scaffolds were evaluated in vitro for cell growth and cytotoxicity with human skin fibroblasts. None of the samples were cytotoxic above the permitted threshold of 50%. This study provides an important increase in performance for the synthesis of poly (L-lactic acid) (PLLA) and further development in the manufacturing of scaffolds which have multiple uses.

Keywords: Poly (L-lactic acid); L-Lactide; Fibrous scaffolds; Cytotoxicity

Introduction

The possible depletion of non-renewable raw materials and the overall increase in pollution has promoted the search for renewable sources for the production of polymers, as well as the synthesis of biodegradable polymers.1,2 Aliphatic polyesters have emerged for short and medium term applications such as sutures, films, dental materials, bags, containers for beverages and disposable materials, due to their biocompatibility and biodegradability.3 Among these, polylactic acid, PLA, has been the most studied, for several reasons: it comes from a natural source, is biodegradable and its degradation products are absorbed by the body.4 It has also been suitable for creating devices such as porous blocks, films and screws. These bioabsorbable devices have been widely used for orthopedic purposes and they also possess great advantages compared with titanium implants or other metal implants because they do not erode the bone when implanted. Additionally, they do not require an extra surgery for removal, which reduces medical costs and allows the gradual recovery of tissue as the material is degraded. Recent advances include fabrication of stents subjected to surface treatments such as pigation to control protein deposition.5 Most of the applications require the production of high molecular weight PLA which can be achieved using either high boiling solvents or depolymerization of oligomers to obtain the dimer called lactide, followed by Ring Opening Polymerization (ROP). This process has been studied with regard to the influence of many parameters such as temperature, pressure, percentage and nature of the catalyst on the molecular weight of the polymer.6 Several methods have been designed for the production of scaffolds, such as membranes or porous blocks, by solvent evaporation, lyophilization, particulate leaching,1,7,8 foams by injection and subsequent leaching of carbon dioxide at high pressure,10 porous fabrics by electrospinning,11,12 phase separation,7,9,13 additive manufacturing,14 laser sintering,15,16 and 3D printing.13,15-16 Electrospinning is a technique patented in 1934, by Formhals17 where a polymer solution is pushed through a needle with the aid of an electric field to produce fibers with a diameter ranging from nanometers to micrometers.

This article describes the synthesis of L-lactide (L-Lc), for subsequent ring opening polymerization (ROP) to PLLA with different molecular weights and electrospinning tests to obtain fibrous scaffolds for in vitro cell growth.

Experimental procedure

Synthesis of low molecular weight PLA (Oligomers)

40mL of L-(+)-lactic acid (LA) (88 to 92%, Scharlau) were placed in a dry 200mL round bottom flask immersed in a sand bath with constant stirring. Starting at 70°C, and 700mbar, temperature and pressure variations were performed every five minutes, continuing with 90°C, 400 mbar, 100°C, 300 mbar, and from 110 to 150°C, 100 mbar. After reaching 150°C, 100 mbar, the process continued until complete water removal to yield 20g of the PLLA oligomer (450-1000g/mol, density 0.825g/mL) following procedures previously established in our laboratory.18

Crude Lc synthesis

The oligomer was distilled at 250°C to 260°C, adding 1.5% SnCl2, 2H2O (Riedel - de Haën ), collecting the crude Lc in a 200mL round bottom flask, dipped in an ice bath, and connected to a condenser with circulating water at 4°C. This procedure supplied 18.74g of crude Lc, with a yield of 55.20%. After recrystallization with ethyl acetate in a 1:2 weight ratio with respect to crude Lc, 15.45g (46.50% yield) were produced.

Synthesis of high molecular weight PLA through ROP

2g of L-Lc, 13.84µL of Sn(Oct)2 (Alfa Aesar), and 220.04mL of 1-octanol (Merck) were weighed in a two-necked 100 mL flask under argon atmosphere in dry bag, under constant stirring at 120rpm. Subsequently, the reaction flask, under argon atmosphere, was introduced into an oil bath and heated to 140 to 145°C. This reaction...
conditions were maintained for 24, 48, 72 and 96 hours respectively, while the obtained PLLA acquired a solid consistency.

**Purification of high molecular weight PLA**

The above process was repeated to obtain 10.0g of PLA, which was dissolved in the least amount of chloroform (Merck), with stirring at room temperature. Then, twice the volume of methanol (Merck) was added to form a milky suspension that, eventually, precipitated. This white solid was filtered in vacuum Buchner funnel to give 9.4g (mp: 92 and 94°C) and finally stored in a dessicator.

**Spectroscopic properties and characterization**

**Thermal analysis:** The L-Lc and PLLA were analyzed by differential scanning calorimetry (DSC) and thermogravimetric (TGA) in a TA-modulated SDTQ600 equipment at a heating rate of 10°C /min under nitrogen atmosphere with a flow rate 100mL/min from 35°C to 350°C, using samples weighing approximately 10mg.

**Molecular weight determination by capillary viscometry:** PLLA molecular weights were determined using a 1C E-999 Ubbelholde viscometer (NOOC-200), chloroform (CHCl₃) as solvent and solutions with concentrations of PLLA 0.02, 0.01, 0.007, 0.005, 0.003, 0.001g/ cm³, in a thermostatic bath at 25°C. Elution times were measured by triplicate for each concentration and the relative viscosity calculated as 1.0279. The specific viscosity was obtained by extrapolating to zero in the plot ηsp/C vs. the square root of the concentration. The molecular weight was obtained by the Mark-Houwink-Sakurada equation (1).

\[
[\eta] = \kappa M^{\alpha}
\]

Where: \( \kappa \) and \( \alpha \) are constants characteristic of each polymer and \([\eta]\) is the intrinsic viscosity. \( \kappa = 6.67\times10^3 \) and \( \alpha = 0.67\)

**Electrospinning of PLLA obtained through ROP**

The parameters to obtain the fibers were: voltage 20kV, distance 14cm; flow rate of 8mL/min. A 40% solution of PLLA (295,000g/mol) in CHCl₃, was prepared and placed in a 5mL syringe attached to a hose, the voltage source (20kV) was connected with a flow rate of 8mL/min. The distance between the needle and the foil was 14cm; flow rate of 8mL/min. A 40% solution of PLLA (295,000g/mol) in CHCl₃, was prepared and placed in a 5mL syringe attached to a hose, the voltage source (20kV) was connected with a flow rate of 8mL/min.

**Preparation of PLLA samples for cell growth:**

These fibrous PLLA scaffolds were used in triplicate for fibroblasts growth, which acts as an extracellular matrix (ECM) where cells may adhere and grow, thus enhancing the development of new tissue, transportation of nutrients and removal of unwanted materials.

**Cytotoxicity test of fibrous scaffolds**

An *in vitro* cytotoxicity test was performed with the LDH Roche® kit. This test is a colorimetric test used to quantify cell lysis and it is based on the LDH measure of the activity released from the cytosol of lysed cells in the supernatant. L929 cell line (ATCC CCL1, NCTC clone 929) is a human skin fibroblast line subcloned from the line L.33

**Preparation of working solutions**

A 1mL of the catalyst aqueous solution was added and stirred for 10 minutes. This solution can be kept for weeks at 2-8°C. The dye solution is stored at 2-8°C. For the 100 trials, 100μL of the catalyst with 11.25mL of dye solution was mixed.

**Cellular culture**

The number of defrosted cells was determined by the tripan blue method and they were seeded in a concentration of 150,000 cells per well in a 24 well (FALCON) plate with a growing medium (Modify medium Eagle’s Dulbeco (DMEM), supplemented with 10% fetal bovine serum (FBS) (GIBCO BDRL), 2mM L-glutamine buffered saline (PBS)). The cell plate was incubated for 48h with CO₂, until a growth of 100%. After incubation time and further verification of cell confluence, fibrous PLLA scaffolds were added as the final product. All triplicate samples were mounted as shown in Figure 1. The 24-well plate containing the samples described above and cells were incubated at 37°C in CO₂ incubator for 24 hours.

**Figure 1** Schematic representation of a 24-well plate. Note that there are three wells on each plate with each of the treatments A, B, C, D, E and F. In addition, arranged in each plate, there were two controls: (1) a high control (CA) and (2) an under control (CB), each in triplicate (three wells).

The arrangement of the materials to be analyzed in 24-well plate was:

- **A:** Cell culture medium (white).
- **B:** Cells+pure PLLA 96h.
- **C:** Cells+commercial PLLA.
- **D:** Cells+pure 80% PLLA 48h and commercial 20% PLLA.
- **E:** Cells+pure 90% PLLA 72h and commercial 10% PLLA.
- **F:** Cells+pure 90% PLLA 96h and commercial 10% PLLA.
- **CA:** Cells+culture medium+triton solution.
- **CB:** Cells+culture medium.

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Cytotoxicity test with LDH Kit ® ROCHE

After the incubation time, 100μL of supernatant were transferred from each well of the first plate to the corresponding wells in a 96 well plate, 100μL of LDH reagent were added and the incubated plate at room temperature was protected from light for 30 minutes. After this time the optical density was determined using a micro-plate reader for ELISA, STAT-FAX 2100, at a wavelength of 456nm with a reference filter 492nm. In order to determine cytotoxicity, the LDH technique was performed with measurements at 72h post incubation with materials and incubation for 30 minutes with the LDH reagent with the Stat Fax-2100 at a wavelength of 456nm with a reference wavelength of 490nm.

Results and discussion

Commercial LA (88-92%) consists of dimers, trimers and Lc.24 Upon heating a PLA oligomer is formed as shown in Figure 2.

![Figure 2](image)

The molecular weight of the formed oligomer can be controlled using different catalysts, functionalizing agents or by changing the polymerization conditions.23 Direct poly-condensation of LA in situ is not applied on an industrial scale because of the competition between the formation of Lc and simultaneous degradation that occurs in the reaction.26,27 The second stage is the de-polymerization of the oligomer to give Lc, as illustrated in Figure 3.

![Figure 3](image)

This molecular weight is adequate for electrospinning of the samples.

Thermal analysis of the synthesized compounds

L-lactide: The melting temperature (T_m) was 95.65°C and the decomposition temperature (T_d) 195.12°C. The values found in the literature differ greatly for example Fuentes et al.36 reported a melting temperature of 119.8°C, a difference that could be due to the contribution of DL-racemic mixture. PLLA: The T_m observed was 70.56°C, T_g 143.6°C and the T_d is 306.43°C. The thermal characteristics of PLLA are important for manufacturing fibrous scaffolds by electrospinning. A relatively high T_g with respect to the reported 70.54°C shows that the synthesized PLLA is high molecular weight. It should be noted that T_g and T_m increase as the molecular weight increases and are strongly influenced by the degree of crystallinity of PLLA. A relatively high T_g of 68°C for a highly crystalline PLLA was reported by Gupta.26 The T_g for 96 h PLLA obtained by us is very close to this temperature, indicating that the obtained PLLA was highly crystalline. It was also found that the PLLA with a T_g of 70.2°C has a 49% degree of crystallinity, which indicates a satisfactory percentage of crystallinity for our 96 h PLLA, that would make it suitable for electrospinning.27

Thermogravimetric Analysis (TGA) of L-Lc: This analysis was performed on the same machine and the same parameters of the experimental part and results session. It is concluded that L-Lc starts...
its decomposition at a temperature of 108.5°C with a weight loss of 6.7% and reach a high degree of decomposition at 195, 1°C with a weight loss of 95%.

**TGA analysis of PLA 96h:** The PLA started decomposition at a temperature of 250.33°C with a rate of 95.03% weight and achieved a high degree of decomposition at 318.3°C with the weight percentage of 1.81%.

### Electrospinning

The parameters used are similar to those reported by Zong et al. A 40% (W/V) of a 90:10 mixture of PLA molecular weight 295,000g/mol and commercial PLA was prepared and subjected to electrospinning and the fibers were collected on aluminum rectangular boxes. These samples were used for cell growth.

### Cytotoxicity tests

To calculate cytotoxicity is necessary to check the three controls in this experiment: the first control is the cell culture medium, which provides information on the activity of LDH. The absorbance value obtained in this control should be subtracted from the values of the other controls. The second control is labeled as “under control”, which provides information on the activity released from untreated normal cells. The third control is labeled as the “high control”, which provides information on the maximum activity of the LDH enzyme released.

Finally, the equation (2) is used to determine the percentage of cytotoxicity of each of the compounds tested.

\[
\text{Cytotoxicity} \% = \frac{\text{Sample Lecture} - \text{CB}}{\text{CA} - \text{CB}} \times 100
\]  

(2)

The percentage cytotoxicity is calculated by applying equation (2) to all samples. Table 1 shows the results of cytotoxicity of PLLA.

#### Table 1 Results of the cytotoxicity test of the PLLA samples

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<tr>
<th>Muestras</th>
<th>Cytotoxicity (%)</th>
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<tr>
<td>PLLA 96h</td>
<td>49.34±0.01</td>
</tr>
<tr>
<td>PLLA comercial</td>
<td>23.00±0.01</td>
</tr>
<tr>
<td>80% PLLA 48h pure y 20% PLLA comercial</td>
<td>42.76±0.01</td>
</tr>
<tr>
<td>90% PLLA 72h pure y 10% PLLA comercial</td>
<td>32.12±0.01</td>
</tr>
<tr>
<td>90% PLLA 96h pure y 10% PLLA comercial</td>
<td>42.93±0.01</td>
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According to Table 1, the synthesized PLLA have the greatest value of all. The commercial PLLA has the lowest value. Nevertheless, Lozano (2013) argues that the fibrous blocks made of PLLA by less than 50% are not considered cytotoxic. Therefore, all PLLA synthesized in this study did not exhibit cytotoxicity and are considered useful in applications with living cells. The next step is to carry out histological tests in vivo.

### Conclusion

We found that mixtures of synthesized PLA (295,000g/mol) with commercial PLA, 90/10 by weight were successfully processed by electrospinning obtaining fibrous matrices that gave favorable results in cytotoxicity tests with human skin fibroblasts.

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### Conflict of interest

The author declares no conflict of interest.

### References


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