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# Advanced thermal analysis and transport properties of stereocomplex polylactide

Nagihan Varol

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Normandie Université

## THESE

Pour obtenir le diplôme de doctorat

Spécialité : Physique

Préparée au sein de l'Université de Rouen Normandie

### Analyse Thermique Avancée et Propriétés de Transport de Matériaux PolyLactide Stéréocomplexe

Présentée et soutenue par  
Nagihan VAROL

Thèse soutenue publiquement le 26 Juin 2019 devant le jury composé de

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### Advanced Thermal Analysis and Transport Properties of Stereocomplex PolyLactide

Presented and defended by  
Nagihan VAROL

Defended on 26 June 2019 in front of the dissertation committee composed of

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### General Introduction

Thermoplastics are currently main polymer materials used in the packaging, automotive, electronics, and medical applications. Due to the environmental context of the sustainable innovations, biobased and biodegradable polymers occupy an increasingly important place in the eco-design of materials [1]. The increase of sustainable concerns of the petroleum-based polymers results in a growing interest in the biopolymers, in particular poly(lactic acid) (PLA)-based materials [2]. Besides, PLA requires 25-55% less energy to be produced than petroleum-based polymers and researches show that this value can be further reduced to less than 10% in the future [3]. In addition, good thermal properties of PLA-based polymers increase the competition with the fossil-based polymers. Nevertheless, the functional properties of PLA-based materials should be improved in terms of their thermal, mechanical and barrier properties in order to correspond to food packaging requirements. For this purpose, many different strategies have been already applied; physical or chemical modifications (crosslinking), blending with other components, and addition of plasticizers or compatibilizers [4-7].

When the interactions between the polymers with the different tacticity predominate over those with the same tacticity, a stereoselective association takes place, which is called stereocomplexation or stereocomplex (sc) formation. Such association allows to form new polymer with enhanced physical properties compared to the parents' polymers. The presence of the chiral carbon in the skeletal chain of PLA yields two stereoregular enantiomers, i.e. poly(L-lactide) (PLLA) and poly(D-lactide) (PDLA). The mixture of homochiral PLLA and PDLA macromolecules leads to the formation of stereocomplex PLA (sc PLA) by the stereocomplexation reaction. Stereocomplexation improves PLA properties through increasing the crystallization ability and crosslinkings between enantiomers. Sc PLA attracts great attention for packaging industry as 2<sup>nd</sup> generation PLA due to the enhanced thermal and mechanical properties. Sc PLA has higher melting temperature ( $T_{m, sc\ PLA} = 240\text{ }^{\circ}\text{C}$ ) compared to the homopolymers (PLLA and PDLA) and co-polymer (PL-DLA) ( $T_{m, PLA} = 180\text{ }^{\circ}\text{C}$ ) [8, 9]. Furthermore, it has been reported that the stereocomplexation enhances the mechanical properties, i.e. increase in ultimate tensile strengths from 31 MPa for pure PLLA to 50 MPa for

sc PLA, and in elongation at break from 3.1 - 5.8% to 15 - 160% depending on the molecular weight of PLAs is observed [10-12].

Unlike racemic mixtures of small chiral molecules, in the case of a polymer, the length of macromolecular chains (molecular molar mass) will strongly influenced the mobility and the structural arrangement of the material. The numerous studies dealing with the sc PLA formation, its microstructure, and degradation have been carried out. Although many researchers focused their attention on the sc crystals species in order to understand the sc PLA properties [13, 14], the behavior of the amorphous phase and the transport properties of sc PLA is still enigmatic topic. Besides it is known that the barrier properties of PLA are still too low in the presence of moisture for the industrial packaging requirements. In order to overcome this problem the addition of lamellar nanofillers [15], the plasma surface treatment [16] or elaboration of polymer blends [17] were proposed in the literature.

In this work, the sc PLA films were elaborated by two methods, i.e. solution casting and extrusion process. Furthermore, the enhancement of the physical and transport properties of PLA-based materials was noted by stereocomplexation, i.e. without the additives, surface treatments, chemical modifications, or inorganic fillers. The transport properties of small molecules ( $H_2O$ ,  $N_2$ ,  $O_2$  and  $CO_2$ ) were linked with the molecular mobility over a wide range of temperatures (from room temperature to the glass transition temperature) in order to understand the role and evolution of the free volume. The aim of this work is to study new PLA-based material obtained from PLLA and PDLA by solution casting and extrusion with enhanced transport properties. The originality of this work bases on the complementarity of two approaches (physics and chemistry) for a better understanding and a broader vision of the properties of the amorphous phase and the evolution of the free volume in stereocomplex polymer which is not studied yet. We studied wholly amorphous and isotropic crystalized materials in order to generate more or less confined amorphous phases. This research work and results have been presented as follows:

- the state-of-art of stereocomplexation and properties of amorphous and semi-crystalline polymer materials is presented in the first chapter. The effective parameters on stereocomplexation are discussed with the elaborations methods.

## *GENERAL INTRODUCTION*

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The properties of the glassy-polymers are presented to exhibit different theories and concepts related to the glass transition phenomenon. This information allows us better understanding the molecular dynamics and relaxation phenomena of the glass-forming liquids and the glassy state;

- the second chapter presents the materials as well as instrumental techniques used in this work, namely calorimetric investigations, microstructure analysis and permeametric analysis;
- the influence of the elaboration methods on the stereocomplexation is developed in the third chapter. The selection procedure of the sc PLA sample elaboration is detailed. The effective parameters, i.e. solvent nature, proportion of homopolymers, elaboration conditions, were compared and discussed;
- the fourth chapter reports glass transition and molecular mobility of the homopolymers and wholly amorphous sc PLA. Fast scanning calorimetry (FSC) and dielectric relaxation spectroscopy (DRS) methods were combined to perform the kinetic study of the molecular dynamic. The aim of this investigation is to observe the influence of the tacticity on the molecular dynamics in the stereocomplex structure;
- the last chapter presents the transport properties of sc PLA in relation to the molecular mobility of the semi-crystalline and amorphous samples. The microstructure of sc PLA is also reported to highlight the structural influences on the barrier properties of sc PLA. Special attention has been given to the water permeability as a function of temperature.

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In this chapter the bibliographical research related to PLA-based materials is presented. The different parameters affecting PLA stereocomplexation, its diverse properties have been summarized. The scientific basics and concepts that can be applied were briefly explained and discussed. Besides the scientific concepts, the history of poly(lactide)-based materials and their importance were presented to define the innovation of this work.



### 1.1 Chirality and Stereoisomers

The concepts of chirality and stereoisomerism offer a great importance to understand the physical and theoretical reasons behind the formation and structures of numerous molecules. A non-superimposable set of two molecules that are mirror image to one another is known as chiral molecules [1] (Fig 1.1). The chiral compounds share the same atomic shape but they are different in structural arrangements; such example of chiral compounds is lactides.

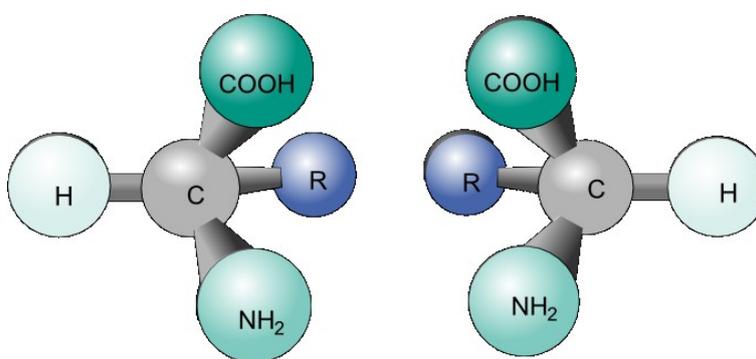


Figure 1. 1. Two enantiomers of a tetrahedral complex [1].

The isomers of chiral compounds are called stereoisomers. In contrast to the well-known constitutional isomerism, which develops isotopic compounds simply by different atomic connectivity, stereoisomerism generally maintains equal atomic connections and orders building blocks as well as has the same number of atoms and elements. Spatial arrangement of atoms concerns how different atomic particles and molecules are situated in the space around the carbon chains. In this sense, the spatial arrangement of a polymer chain will be different if one of the atoms in the polymer chain is shifted even by one degree in any three-dimensional directions. This opens up a broad possibility formation of different molecules, each with their unique placement of atoms in the three-dimensional space [2].

### 1.2 Lactides

Lactides are produced by the depolymerization of poly(lactides) under reduced pressure to give a mixture of *L*-lactide, *D*-lactide and/or *meso*-lactide isomers (Fig 1.2) [3]. The different percentages of the obtained isomers depends on the lactic acid isomer feedstock, temperature and catalyst.

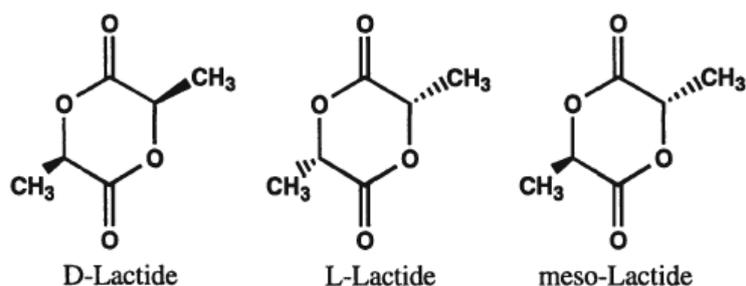


Figure 1. 2. Chemical structures of *L*-lactide, *D*-lactide and *meso*- or *LD*-lactide isomers [3].

### 1.3 Poly(lactide)

Due to environmental concern, the development of biosourced and biodegradable packaging materials has attracted great interest to replace petroleum-based plastic packaging materials. PLA is one of the mostly used biodegradable thermoplastic aliphatic polyester derived from renewable bio-resources [4, 5]. It is used in many applications from biomedical devices and tissue engineering to packaging due to its excellent biocompatibility and sustainability [6, 7]. In particular, PLA is used in food packaging application due to its numerous advantages:

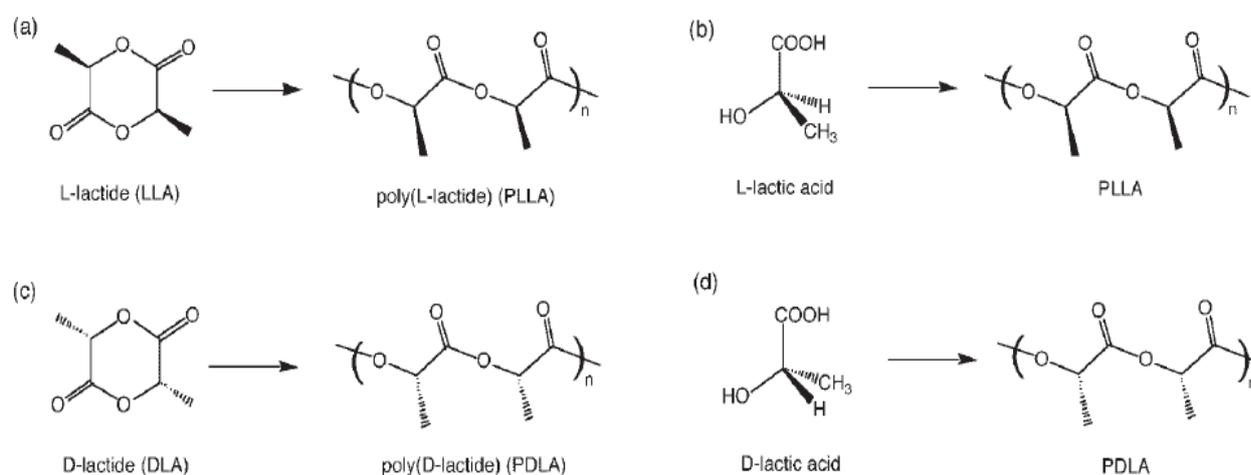
- it can be obtained from renewable bio-sources
- it provides significant energy savings
- it is recyclable and compostable
- it is useful to improve farm economies
- the physical and mechanical properties can be tunable through the polymer architecture, i.e. grafted homopolymers by graphene oxide [8] and star-shaped isomers [9].

Many studies showed that PLA is an economically feasible material to use as a packaging polymer. However, PLA is a very brittle material and the toughness in its raw state is insufficient to resist a high level of impact strength [10]. The thermal and mechanical stability of PLA limits its wide applications that still need higher stress levels and higher barrier properties. In addition, the low crystallization rate of PLA causes low heat distortion temperatures (HDT) [7, 11]. Therefore, there are still so many challenges for the full applications in utilizing neat PLA-based materials. Many solutions have been developed to improve the property deficiency of PLA, including plasticization, copolymerization and melt

blending with flexible polymers [12]. Such modifications enhance PLA's elongation at break and reduce brittleness. However, the stiffness of the blended materials is also an important property for the applications. Therefore, the new production strategy is developed without compromising the stiffness and tensile strength to enhance the thermal and mechanical stability of neat PLA in a broad temperature range.

### 1.3.1 Chain structure and production of PLA

PLA is the simplest hydroxyl acid with a chiral (or an asymmetric) carbon atom and it exists in two optically active configurations, the *L*- (+) and *D*- (-) isomers which are presented in Fig 1.3. The basic configurational unit of PLA can be manufactured by carbohydrate fermentation (as shown in Fig 1.4) or chemical synthesis (as shown in Fig 1.5).



**Figure 1. 3.** Synthesis and molecular structures of PLLA [(a) and (b)] and PDLA [(c) and (d)] [13].

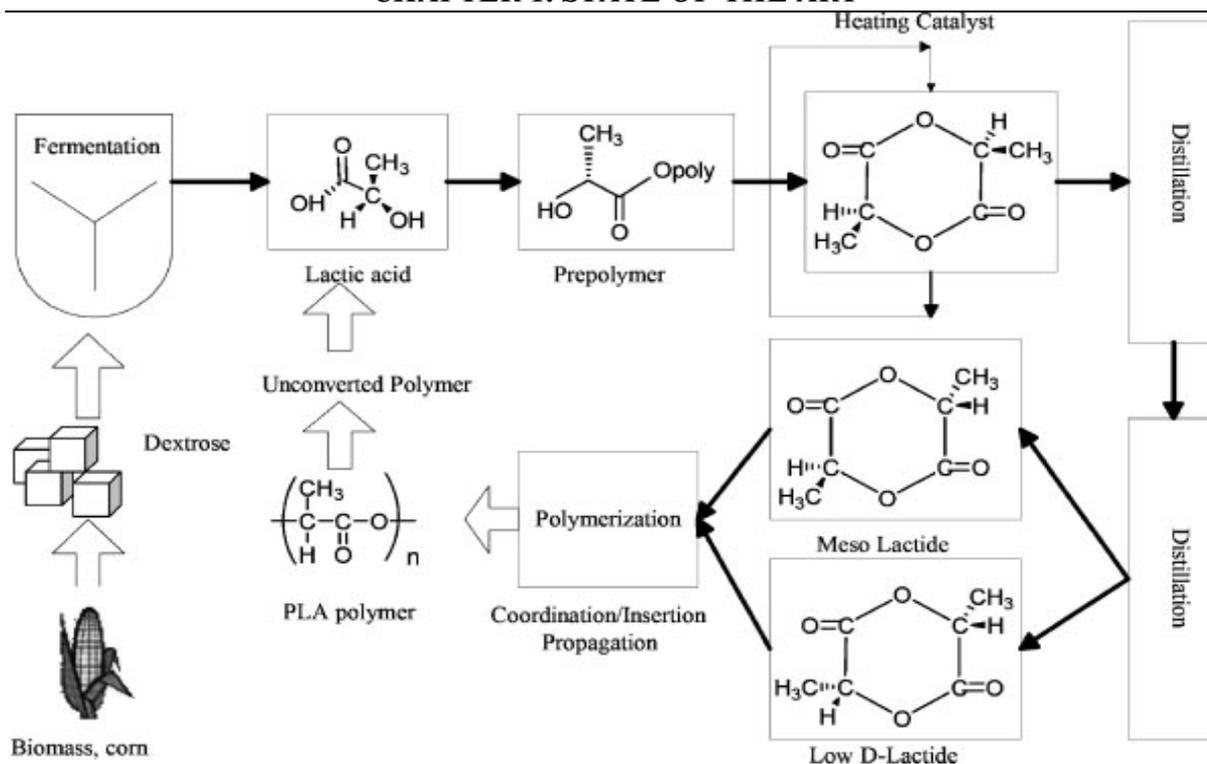


Figure 1. 4. Schematic illustration of the production of PLA, image was adopted from Gruber et al. [14].

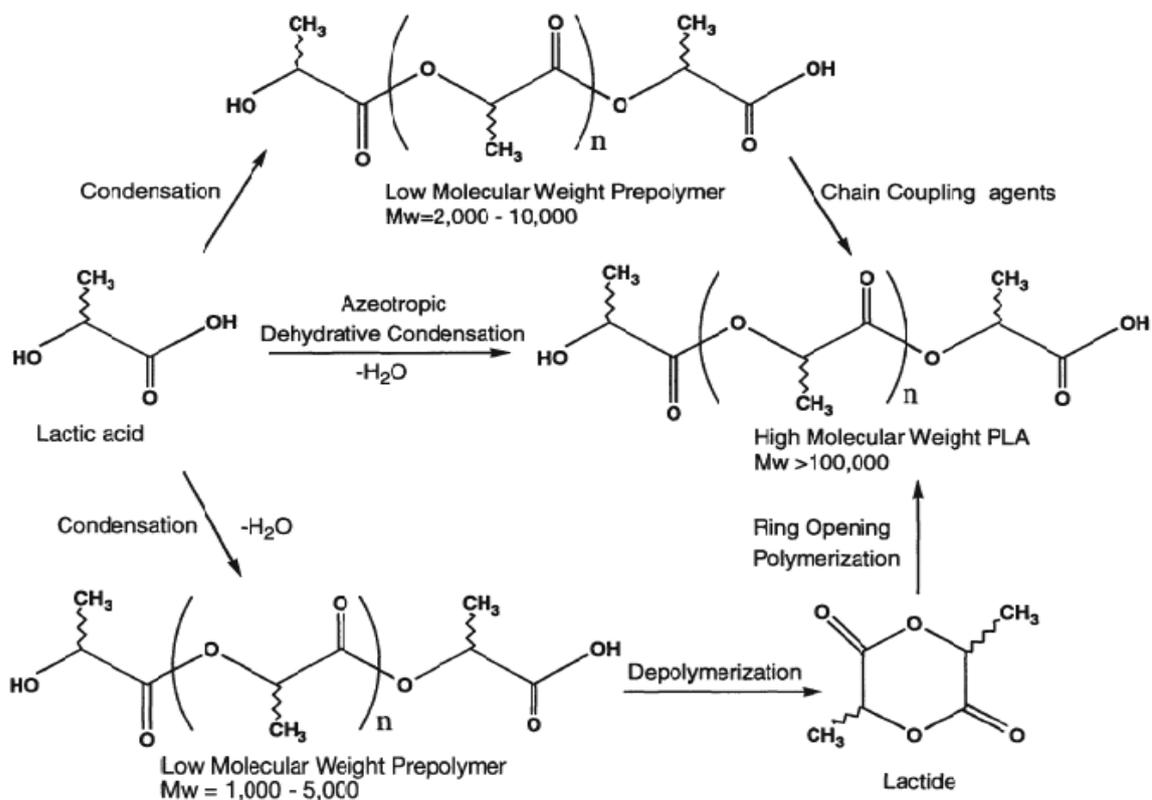


Figure 1. 5. Schematic illustration of chemical synthesis of low and high molecular weight PLA, image was adopted from Hartmann et al.[3].

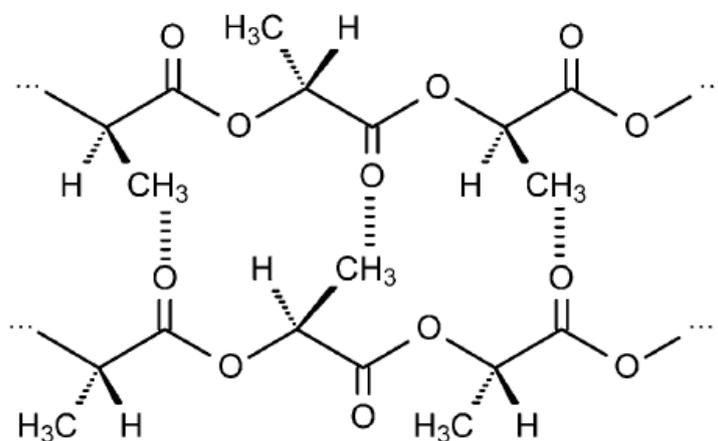
### **1.3.2 Crystal structures of PLA**

Different crystal structures of PLA have been reported according to the crystal formation and the crystallization conditions. The well-known  $\alpha$  crystals obtained in conventional melt and solution crystallization conditions were first reported by De Santis and Kovacs [15], then further investigated in a numerous studies [16-18]. Zhang et al. [19] noted based on Wide Angle X-ray Diffraction (WAXD) and Infrared (IR) spectroscopy, the slightly different  $\alpha'$  crystals of PLA crystalized below 120 °C. The chain conformation and crystal structure of  $\alpha'$  form is similar to  $\alpha$  form, but  $\alpha'$  form has a less ordered chain packing. It is proposed that pure  $\alpha'$  crystals are formed at crystallization temperatures below 100 °C, whereas crystallization between 100 °C and 120 °C leads to the coexistence of  $\alpha'$  and  $\alpha$  crystal structure [20, 21]. A  $\beta$  form, firstly reported by Eling et al. [22], is created by stretching the  $\alpha$  form at high draw-ratio and high temperature, i.e. during hot-drawing of melt or solution spun fibers [22, 23]. Melting temperature of the  $\beta$  crystals is about 10 °C lower compared to the  $\alpha$  crystal, attesting that  $\beta$  form is thermally less stable [23]. Then, Puiggali et al. [18] suggested that  $\beta$  form crystal is a frustrated structure with a trigonal cell containing three chains which are randomly oriented up and down. A more ordered crystal formation called  $\gamma$  was also reported by Cartier et al. [24]. In the  $\gamma$  form, which was obtained by epitaxial crystallization of PLA on hexamethyl benzene, two chains are oriented antiparallel in the crystal cell.

### **1.4 Stereocomplexation**

Besides the homocrystallization of PLLA and PDLA, these two enantiomeric chains can co-crystallize together and form a stereocomplex crystals [25]. In contrast to PLLA or PDLA homocrystals, the stereocomplex crystal cell contains one PLLA and one PDLA chain. Well-known examples of stereocomplexation for enantiomeric polymers and for polymers with different tacticity were reported by Pauling and Corey in 1953 [26] and by Fox et al. in 1958 [27], respectively. Since a novel stereocomplex interaction between two chiral polymers was discovered, a third group of polymer complexes are obtained based on hydrophobic or van der Waals interactions. Stereocomplex formation between enantiomeric polymers leads to achieve better physical and mechanical properties than those of pure components.

Interestingly, melting temperature of the stereocomplex is about 50 °C higher than that of PLA homocrystals. In virtue of the different tacticity of PLA isomers, i.e. PLLA and PDLA, the improved interactions between the isomers with different tacticity are governed by stereocomplexation. Due to the different tacticities, a stereoselective association takes place through the stereocomplexation reaction. The presence of chiral carbon in the skeletal chain of PLA yields two stereoregular enantiomers (PLLA and PDLA) that allows forming the stereocomplex PLA (sc PLA) by stereocomplexation, as depicted in Fig 1.6. Stereocomplexation can be obtained by the attracting complementary interactions result from a hydrogen bonding [28]. *L*-lactide and *D*-lactide units can maintain the two helical chains with the opposite configuration together through the multicenter hydrogen bonding interactions, between  $-\text{CH}_3\cdots\text{O}=\text{C}<$  and  $\equiv\text{CH}\cdots\text{O}=\text{C}<$  [29-31]. These improvements arise from a peculiar strong interaction between *L*- and *D*-lactide enantiomers, thus opening a new way of preparing innovative materials.



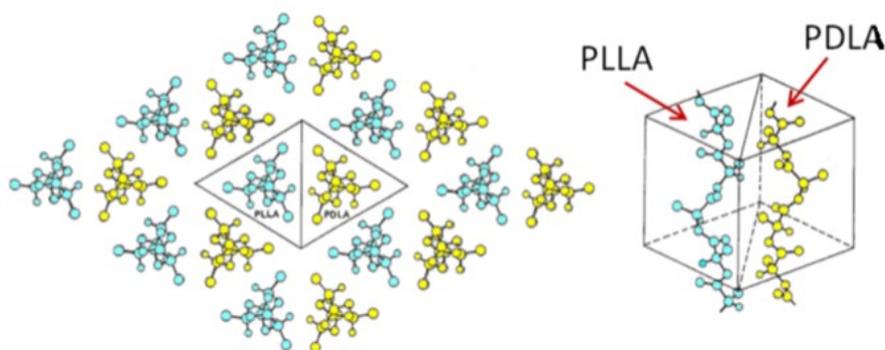
**Figure 1. 6.** Chain structure of sc PLA obtained from the linear poly(*L*-lactide) and poly(*D*-lactide) [32].

## 1.5 Stereocomplex PLA: structure and detection

### 1.5.1 Structure of stereocomplex PLA

Since sc PLA was discovered, Ikada et al. [25] noted that sc PLA has different X-ray diffraction pattern from homopolymers crystals; however, the stereocomplex crystal structure was not determined quantitatively. De Santis and Kovacs [15] reported a right-handed helical conformation of PDLA chain, structurally based on the left-handed helical

conformation of PLLA chain; therefore, they proposed van der Waals forces such as dipole-dipole interactions as the origin of the stereocomplexation (as shown in Fig 1.7).



**Figure 1. 7.** Schematic illustration of the crystal structure of stereocomplex PLA [13].

Based on X-ray measurements and energy calculations, the crystalline structure of sc PLA was investigated in details. The stereocomplex crystals were discovered to be triclinic and trigonal unit cells with dimensions of  $a = 0.916$  and  $1.50$  nm,  $b = 0.916$  and  $1.50$  nm, and  $c = 0.870$  and  $0.87$  nm, with the angles  $\alpha = 109.2^\circ$  and  $90^\circ$ ,  $\beta = 109.2^\circ$  and  $90^\circ$ , and  $\gamma = 109.8^\circ$  and  $120^\circ$ , respectively [33, 34]. Properties of the different types of PLA crystal form are summarized in Table 1.1. The values of density are calculated based on the reported cell parameters and the number of monomers in each unit cell.

Crystal Type	Crystal system	Chain conformation	Cell parameters						$\rho^{\text{theoretical}}$ (g/cm <sup>3</sup> )
			a (nm)	b (nm)	c (nm)	$\alpha$ (°)	$\beta$ (°)	$\gamma$ (°)	
$\alpha$ [15]	Pseudo-orthorhombic	$10_3$ helical	1.07	0.65	2.78	90	90	90	1.25
$\alpha$ [17]	Orthorhombic	$10_3$ helical	1.05	0.61	2.88	90	90	90	1.30
$\beta$ [23]	Orthorhombic	$3_1$ helical	1.03	1.82	0.90	90	90	90	1.28
$\beta$ [18]	Trigonal	$3_1$ helical	1.05	1.05	0.88	90	90	120	1.28
$\gamma$ [24]	Orthorhombic	$3_1$ helical	0.99	0.63	0.88	90	90	90	1.31
sc [33]	Triclinic	$3_1$ helical	0.92	0.92	0.87	109.2	109.2	109.8	1.27
sc [34]	Trigonal	$3_1$ helical	1.50	1.50	0.87	90	90	120	1.27

**Table 1. 1.** Crystallographic parameters of different types of PLA crystals.

In addition, Cartier et al. [34] and Tsuji et al. [35] suggested that chains' conformation is  $3_1$  helix, and each unit cell involves a PLLA and a PDLA chain with the same number of *L*- and

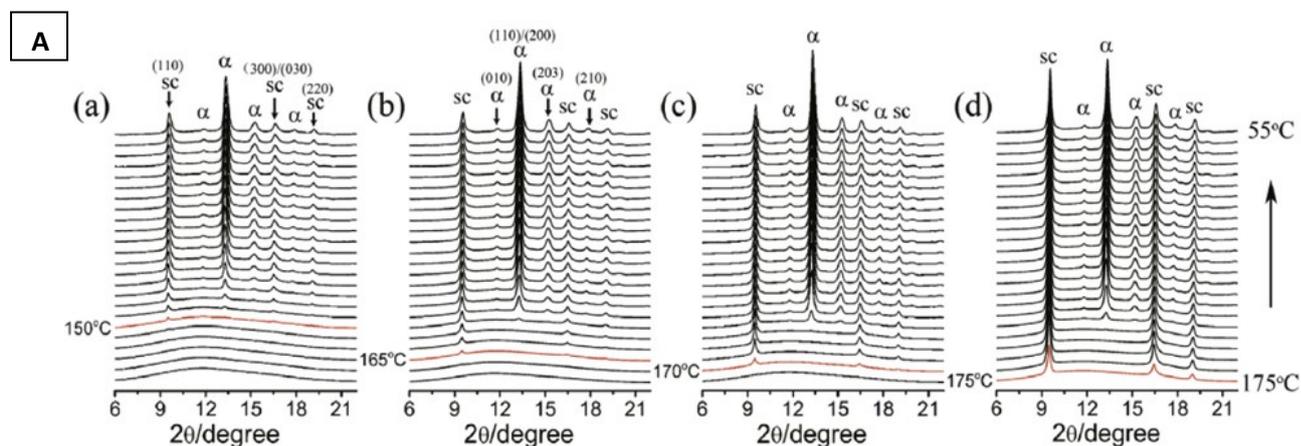
D- units. Therefore, stereocomplex crystals have more extended chains compared to PLLA  $\alpha$  crystals, in which chains are composed of  $10_3$  helices.

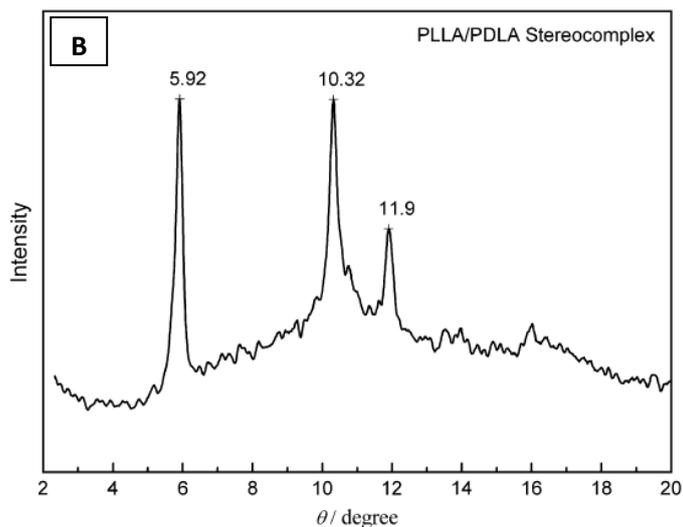
### 1.5.2 Detection of stereocomplex PLA

Sc PLA was detected mostly by X-ray diffraction and DSC analysis due to the different patterns and melting temperature from the homopolymers; thus the presence and fraction of sc crystals in the obtained new material can be easily evaluated.

#### 1.5.2.1 X-ray diffraction

Since sc PLA has different XRD pattern compared to homopolymers, it can be detected without altering the thermal history of polymers. He et al. [36] and Ikada et al. [25] reported diffraction peaks for sc PLA appearing at  $2\theta$  around 12, 21 and  $24^\circ$  while PLA homocrystals diffraction peaks appear at  $2\theta$  equal to 15, 16, 18.5 and  $22.5^\circ$  (Fig 1.8). The quantitative analysis can also be done to evaluate the amount of stereocomplex present in the blends by XRD [35]. Wide Angle X-ray Scattering (WAXS) intensity curves (Fig 1.8A) of PLLA/PDLA blends were obtained by cooled from  $175^\circ\text{C}$  by  $5^\circ\text{C}/\text{min}$  with different shear rates (a)  $0\text{s}^{-1}$  (b)  $50\text{s}^{-1}$  (c)  $100\text{s}^{-1}$  (d)  $178\text{s}^{-1}$  with the temperature interval of  $5^\circ\text{C}$ .

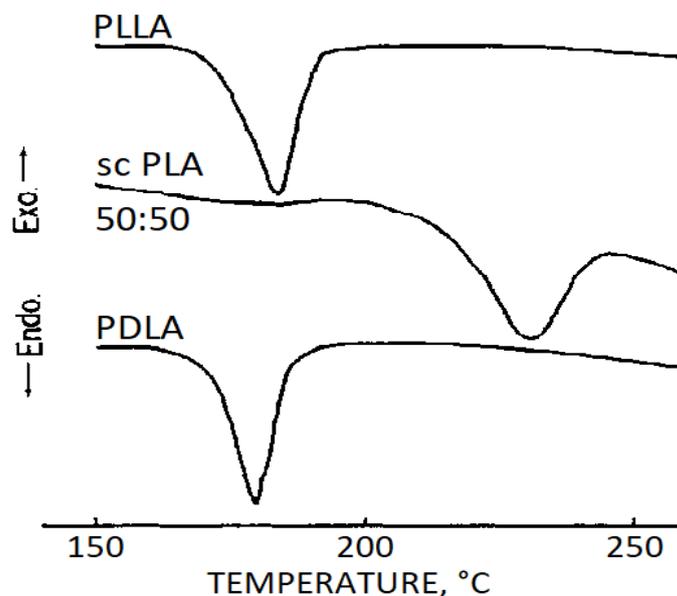




**Figure 1. 8.** Typical (A) WAXS [37] and (B) XRD spectra [36] of PLLA ( $\alpha$ ) and sc PLA (sc) registered with the radiation wavelength  $\lambda = 1.24$  and  $1.54 \text{ \AA}$ , respectively.

#### 1.5.2.2 DSC analysis

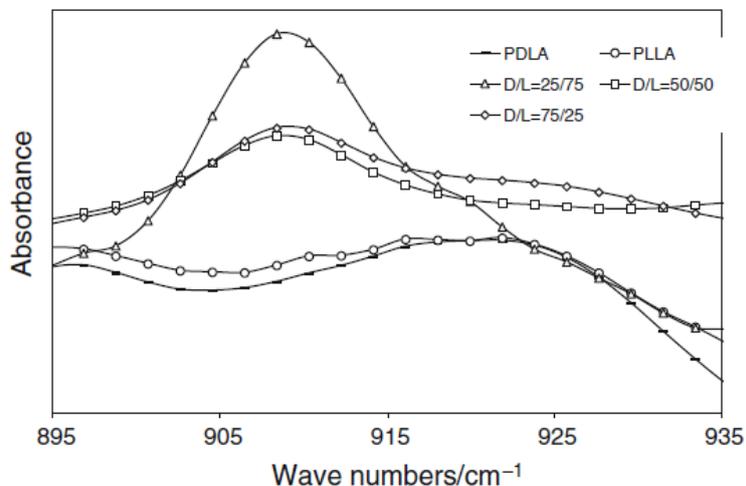
The blends of PLLA and PDLA can form stereocomplex crystals which have a melting temperature about  $50 \text{ }^\circ\text{C}$  higher than those of PLLA and PDLA homocrystals [25]. Due to the higher melting and crystallization temperature, the existence and amount of stereocomplex crystals in the blends can be simply evaluated by DSC analysis (Fig 1.9).



**Figure 1. 9.** Typical DSC curves of homopolymers and sc PLA [25].

### 1.5.2.3 Other techniques

The other methods have also been proposed to detect the presence of stereocomplex in the blends. Such techniques include microscopies and spectroscopies. The former (i.e. scanning electron microscopy (SEM), polarized optical microscopy (POM), transmission electron microscopy (TEM) and atomic force microscopy (AFM)) are generally used to reveal the information about surface morphology, crystals formation as well as crystallization kinetics; while the latter (i.e. Fourier-transform infrared spectroscopy (FTIR), Raman spectroscopy and nuclear magnetic resonance spectroscopy (NMR)) reveal information about the hydrogen bonding in stereocomplex polymers. For instance, there is a new absorption peak at  $908\text{ cm}^{-1}$  in the FTIR spectra of sc PLA due to the hydrogen bonding as compared to homopolymers as shown in Fig 1.10 [38]. The hydrogen bonds undergo rearrangement between  $\text{CH}_3$  group and  $\text{C}=\text{O}$  group and cause a conformational adjustment in the C-O-C backbone (shown in Fig 1.6). In addition, the  $10_3$  helix chain conformation ( $\alpha$  crystals) at  $923\text{ cm}^{-1}$  disappears after stereocomplexation.



**Figure 1. 10.** FTIR spectra of homopolymers and sc PLA (e.g. D/L=25/75 corresponds to 25:75 (w/w) ratio of PDLA and PLLA) [38].

## 1.6 Influence of the different parameters on stereocomplexation

In the racemic mixture of PLLA and PDLA, there is a competition between homocrystallization and stereocomplexation. Several parameters that could influence this competition are follows [25, 33, 37, 39-47]:

- ♦ blending ratio of two homopolymers, PLLA and PDLA [25, 48-50]
- ♦ molecular weight of two homopolymers [49-51]
- ♦ optical purity of the two isomeric polymers [41, 52]
- ♦ temperature and time after blending of two homopolymers in solutions [42, 49, 50, 52]
- ♦ the solvents utilized for polymer casting [49, 51]
- ♦ the structure of co-monomer units and length of the lactide unit sequences in copolymers [53-57].

The most common conditions of blending to form stereocomplex crystallites without homocrystallites include: equimolar mixture of low *MW* *L*- and *D*-lactide homopolymers [25, 42, 49-52].

### 1.6.1 Blending ratio of homopolymers

Ikada et al. [25] has reported that the optimum blending ratio for homopolymers is 50:50 at which only stereocomplex is formed. When deviating from equimolar ratio of PLLA and PDLA, the formation of homocrystals is obvious through the appearance of melting peak at 180 °C. As for the equimolar blend, there is only melting of sc-crystals at 230 °C, as shown in Fig 1.11.

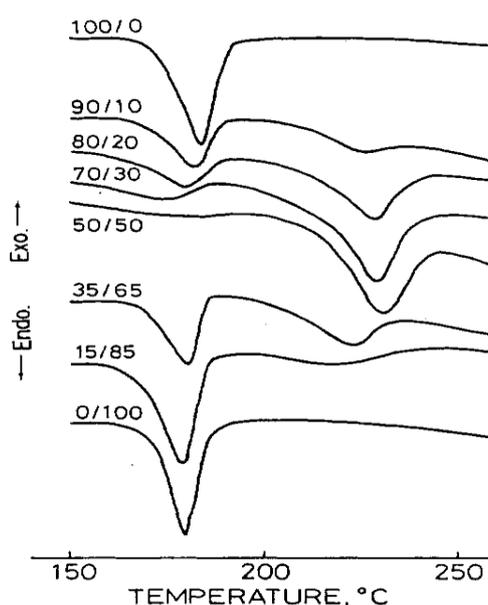
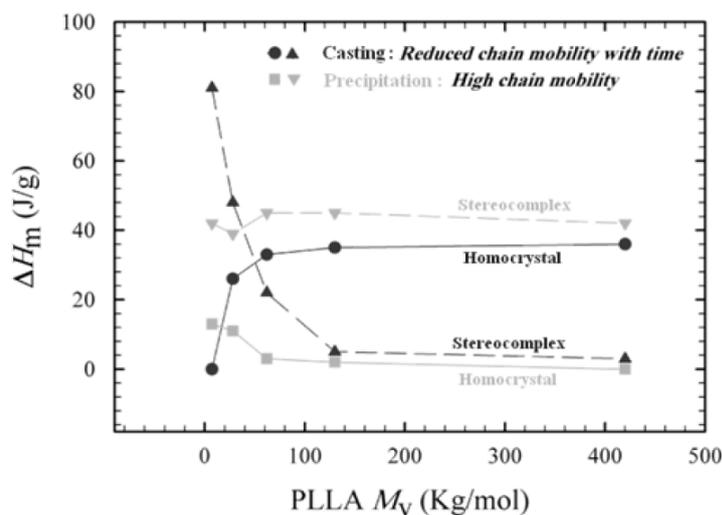


Figure 1. 11. Melting thermograms of polymer blends of PLLA and PDLA in different ratio [25].

In addition, Tsuji et al. [51, 58] indicated that the melting peaks of both homocrystals and stereocomplex were observed when PDLA fraction ( $X_D$ ) was between 0.1 to 0.3 and 0.7 to 0.9, while the pure stereocomplex was detected for  $X_D$  between 0.4 and 0.6 with maximum of the melting enthalpy ( $\Delta H_m$ ) at  $X_D = 0.5$ . The only homocrystallization occurred near  $X_D = 0$  and 1. According to the obtained stereocomplex crystalline structure in which PLLA and PDLA chains are paired in each units with the equilateral triangles structure, it is reported that equimolar blending ratio is the optimum condition for yielding formation of stereocomplex crystals.

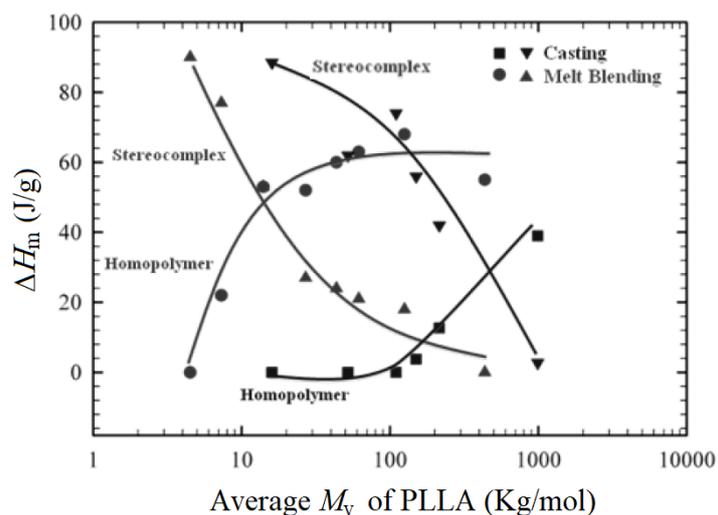
### **1.6.2 Molecular weight of homopolymers**

The work of Ikada et al. [51, 58] and Tsuji et al. [42, 48] demonstrated that the molecular weight of isomers affects the extent of homocrystallization and stereocomplex formation. In their work, not only the impact of the molecular weight of homopolymers was investigated but also different elaboration methods were compared in order to understand the competition between stereocomplex and homocrystals formation. The investigation was carried out by using different viscosity average molecular weight ( $M_v$ ) PLLA and fixed PDLA in a 50:50 ratio of PLLA:PDLA. As shown in Fig 1.12, stereocomplex crystals were obtained over a wide range of  $M_v$ . Stereocomplexation was preferentially obtained when low molecular weight polymer (approximately below 40 kDa) was used in the film casting process. With increasing molecular weight, the formation of stereocomplex decreases while the homocrystals formation increases. When the polymer was prepared by precipitation accompanied by stirring, stereocomplex crystals formation was obtained regardless molecular weight variations, as shown in Fig 1.12. It can be deduced that the chain mobility in the samples prepared by precipitation is higher compared to the samples prepared by solution casting. However, if the precipitation was carried out in high concentration (5g/dL) and/or without stirring, the stereocomplex crystals failed to form. Instead, only homocrystals were obtained [51, 58].



**Figure 1. 12.** Competitive stereocomplexation and homocrystallization in solution casting and precipitation methods [51, 58].

Fig 1.13 shows the effect of blending method as the function of  $M_v$ . Tsuji and Ikada [42] revealed that the stereocomplexation occurred rapidly (e.g. from 90 J/g in 10 kDa polymer prepared by solution casting to 70 J/g when the  $M_v$  is approaching 100 kDa) while homocrystallization required longer induction time (e.g. the  $\Delta H_m$  increases only after 100 kDa).



**Figure 1. 13.** Competitive stereocomplexation and homocrystallization in solution casting and melt blending as the function of  $M_v$  of homopolymer isomers [42, 48].

### 1.6.3 Chain structure of the components

The influence of chain structure on the homocrystallization and stereocomplex formation was investigated by comparing linear and branched PLAs in the racemic mixtures [39]. It was reported that the enhanced nucleation process occurred in the blends due to the branched chain structure. The impact of branched structure was confirmed by reducing cold crystallization temperature (from 140 °C in linear structure down to 111 °C), increasing spherulite density, different growth rate of branched PLAs and different internal morphology during stereocomplexation. In addition, Tsuji and Ikada [59] reported the influence of optical purity on the stereocomplexation of PLLA and PDLA (50:50 ratio) prepared by solution casting. POM micrographs demonstrated that no spherulitic structure was observed for non-blended polymers with optical purity less than 76%. The spherulite density increases with decreasing isothermal crystallization temperature (from 0.2 number/mm<sup>2</sup> at  $T_c = 140$  °C to 1200 number/mm<sup>2</sup> at  $T_c = 100$  °C in the same 98% optical purity PLLA) as well as at higher optical purity (i.e. from 36 number/mm<sup>2</sup> in 76% optical purity PLLA to 1200 number/mm<sup>2</sup> in 98% optical purity PLLA measured at  $T_c = 100$  °C). Furthermore, blended films had higher spherulite density (4200-9700 number/mm<sup>2</sup> at  $T_c = 100$  °C) at any optical purity and crystallization temperature compare to non-blended ones. DSC results depicted that increasing optical purity reduced the cold crystallization temperature and increased melting temperature. No homocrystallization and melting peaks were observed for optical purity less than 90%, and a melting peak corresponded to the stereocomplex crystal was monitored around 230 °C.

Brochu et al. [52] investigated the impact of composition and overall enantiomeric excess on the morphology of sc PLA. The samples were elaborated by solution casting and slow solvent evaporation. The  $\Delta H_m$  of stereocomplex crystals increases with increasing PDLA content, with the maximum value at 50:50 ratio of 100% pure PLLA and 100% pure PDLA isothermally crystallized at 104 °C. Such result can be explained by the prohibition of homocrystals formation in the presence of stereocomplexation. For the blend films that contain 100% pure PLLA and 80% pure PDLA isothermally crystallized at 115 °C, the maximum  $\Delta H_m$  for the stereocomplex crystals was achieved at 35:65 (PLLA:PDLA) blend ratio; while for homocrystals can be observed up to 30:70 ratio of PLLA:PDLA. Furthermore, the amount of homocrystals in the blend of 100% and 80% pure PLLA and PDLA, respectively, is similar to the

blend that contains entirely 100% pure homopolymers. On the contrary, the amount of stereocomplex crystals in the blend films containing 100% pure PLLA and 80% pure PDLA was less than that in the blends with 100% pure isomers. Such a consequence can be attributed to two facts: (1) a decline in the stereocomplexation rates due to the *L*- units in 80% optically pure PDLA chain, and (2) higher molecular weight of 80% optically pure PDLA samples as compared to the 100% pure isomers (higher molecular weight ensures slower chain diffusion to form stereocomplex crystals).

### **1.7 Barrier properties of PLA-based material**

Transport property is one of the most important parameters in food packaging industry. The mostly used polymers should be in the form of non-porous films with high barrier property and selectivity. Permeants (H<sub>2</sub>O and gases: N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>) pass through the polymer films when there is a pressure or activity difference between the opposite film interfaces. Permeation through non-porous polymer films is described by “solution-diffusion” mechanism [60-62]. Based on this mechanism, permeation is a complex process involving different sequence of events:

- 1) absorption of the permeant into the film at its interface applied to the higher pressure or activity
- 2) molecular diffusion of the permeant in and through the film
- 3) desorption and release of the permeant from the film at the opposite interface exposed to the lower pressure or activity.

Due to environmental concerns, the transport properties of traditional PLAs were widely investigated for packaging applications [63-68]. However to our best knowledge, the barrier properties of sc PLA is not well investigated. Tsuji and Tsuruno [69] proposed the influence of crystallinity degree and crystal types on the water vapor permeability (WVP) of PLAs. They prepared pure PLLA and PDLA films with homocrystals and PLLA/PDLA blends with only stereocomplex crystals as the crystalline species. Isothermal crystallization process was applied to the elaborated films in order to obtain films with different crystallinity degree and pure crystals. It was shown that the WVP of studied polymers decreases (barrier properties improves) with increasing degree of crystallinity. Water vapor transmission of homopolymers

became slower (permeability coefficient  $P$  decreased from 2.29 to 1.15 Barrer) with prolong isothermal crystallization time at 140 °C. In the case of sc PLA, the water vapor transmission is faster ( $P$  increases from 1.00 to 1.81 Barrer) when the sample is isothermally crystallized at higher  $T_c$  (ranging from 205 °C to 255 °C). To sum up, the  $P$  value of sc films is 14-23% lower than that of pure PLLA and PDLA films depending on the degree of crystallinity (Fig 1.14).

Nevertheless, there is no sufficient information about the impact of stereocomplexation on the barrier properties of PLA-based materials, especially towards water and gases ( $N_2$ ,  $O_2$  and  $CO_2$ ). In literature, elaborated PLLA/PDLA films were thermally treated after elaboration to increase the purity of crystalline species and degree of crystallinity. Therefore, during this thesis the barrier properties of sc PLA film are studied (cf. Chapter 5). It has been widely accepted that the permeants transport through the “free volume” in amorphous phase disregard of its crystalline structure [70]. Therefore, the next section focuses on the amorphous phase through glass transition concept.

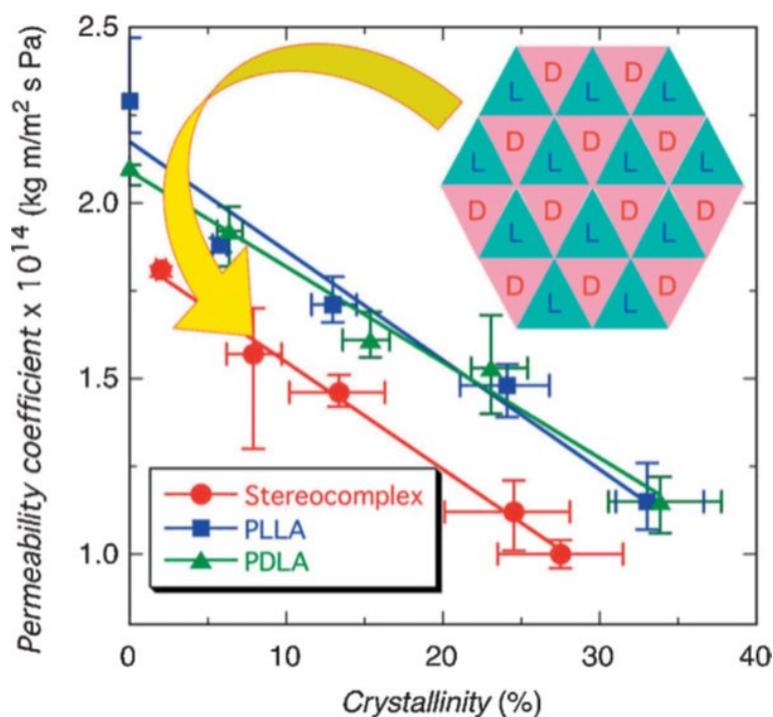


Figure 1. 14. The water vapor permeability coefficient ( $P$ ) dependence on crystallinity degree [69].

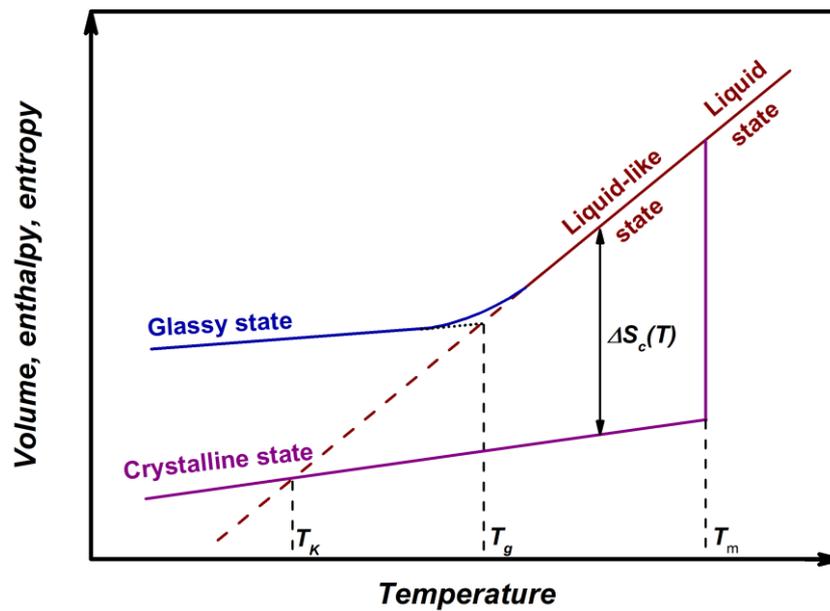
## 1.8 Theory of the glass transition and its phenomenological aspects

The concept of the glass formation and the glass transition temperature ( $T_g$ ) is detailed with theoretical models to describe the molecular mobility related to the glass transition.

### 1.8.1 Glasses and the concept of $T_g$

Glasses are defined as disordered materials, including all solids having a non-crystalline atomic-scale structure. The glasses indicate a transition when heated up to the liquid state. A liquid can form glasses in numerous ways if the applied cooling rate is enough to prevent crystallization of the material [71]. Glasses include different materials display different chemical nature; i.e. oxides [72], chalcogenides [73-75], ionic liquids [76, 77], and polymers [78-80]. The glass transition is a transition from metastable equilibrium (liquid-like state) to the glassy state. Meanwhile glass-forming liquid is cooled down rapidly, its viscosity  $\eta$  increases and reaches a value approximately  $10^{13}$  Poise at a relaxation time close to 100s [81]. The temperature at which the viscosity is reached, is called  $T_g$ . The dynamic phenomenon associated to the  $T_g$ , so called the  $\alpha$ -relaxation, has been widely used to investigate different polymer glass-forming liquids. Most commonly used theories and models are free-volume approach [82, 83], thermodynamic models [84], Adam and Gibbs theory [85], mode coupling theory [86], and coupling model [87]. The glass transition affects different dynamic quantities and thermodynamic parameters such as the viscosity, the heat capacity and the relaxation time which is the required time to recover equilibrium after an external perturbation [88]. Therefore, the  $T_g$  can be measured by different experimental techniques, such as dilatometry measurements and differential scanning calorimetry (DSC) measurements. In Fig 1.15 a typical curve shows the evolution of volume, the enthalpy and entropy of a glass-forming liquids as a function of temperature. Once the melting temperature  $T_m$  is reached, the thermodynamic properties of crystals turn into liquid state species. The molecules are extremely mobile at and above the equilibrium temperature. If the glass-forming liquid is able to crystallize, the thermodynamic properties of the crystalline state can quietly increase towards to higher temperature (path of the purple line). In this case, the entropy of the equilibrium liquid ( $S_{liquid}$ ) would be equal to the entropy of the crystal ( $S_{crystal}$ ) at a temperature, so called Kauzmann's temperature ( $T_K$ ).  $T_K$  is inaccessible and can only be determined by extrapolation, so this situation is referred to the Kauzmann's paradox [89-91].

If the liquids were cooled down by sufficient cooling rate to avoid crystallization, the system will reach in a metastable state, so called the liquid-like state (cooling path on wine-red line). Meanwhile the temperature decreases along with the mobility of the molecules to reach the temperature where the liquid state seems to be frozen, the  $T_g$  is reached resulting in a discontinuity in the evaluation of the thermodynamic properties. Therefore, the  $T_g$  is also commonly defined as the intersection of the liquid and the glassy lines of the thermodynamic properties [92, 93].



**Figure 1. 15.** Schematic representation of the thermodynamic properties of a glass-forming liquid as a function of temperature. Wine-red dotted line corresponds to the thermodynamic equilibrium.  $T_K$ : Kauzmann's temperature,  $T_g$ : glass transition temperature,  $T_m$ : melting temperature,  $\Delta S_c$ : configurational entropy  $\Delta S_c = S_{\text{liquid}} - S_{\text{crystal}}$ .

The glass transition kinetic is demonstrated by the dependence of the  $T_g$  on the cooling rate [94-96]. The faster cooling rate the higher  $T_g$  value and the slower cooling rate the lower  $T_g$  value, as depicted in Fig 1.16. Thus, the  $T_g$  is not intrinsic temperature because of the dependency on the cooling rate. The cooling rate is defined by the first derivative of the temperature  $T$  by time  $t$ :

$$\beta_c = \frac{dT}{dt} \quad (1.1)$$

By estimating the cooling as the series of temperature steps, the eq 1.1 can be expressed as the time  $t$  taken towards each temperature  $T$ :

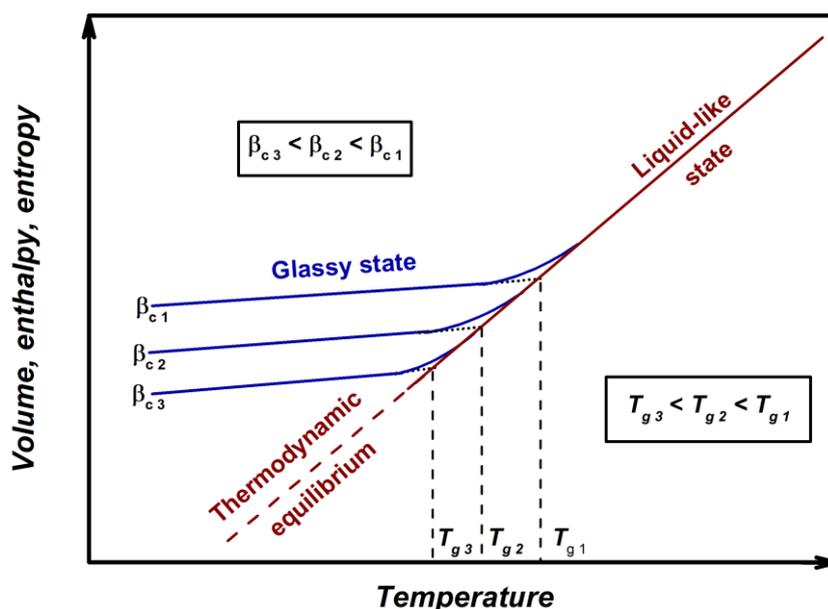
$$\Delta t = \frac{\Delta T}{\beta_c} \quad (1.2)$$

Hereby, higher cooling rate resulting in a shorter time between each  $T$ . Such result allows us to define the idea of relaxation time  $\tau$ , which is required time by the system to reach the equilibrium after external perturbations. Therefore, the liquid falls out equilibrium when  $\Delta t$  at each  $T$  is smaller than  $\tau$ .

$\Delta t \gg \tau \rightarrow$  liquid and liquid like state

$\Delta t \approx \tau \rightarrow$  glass transition

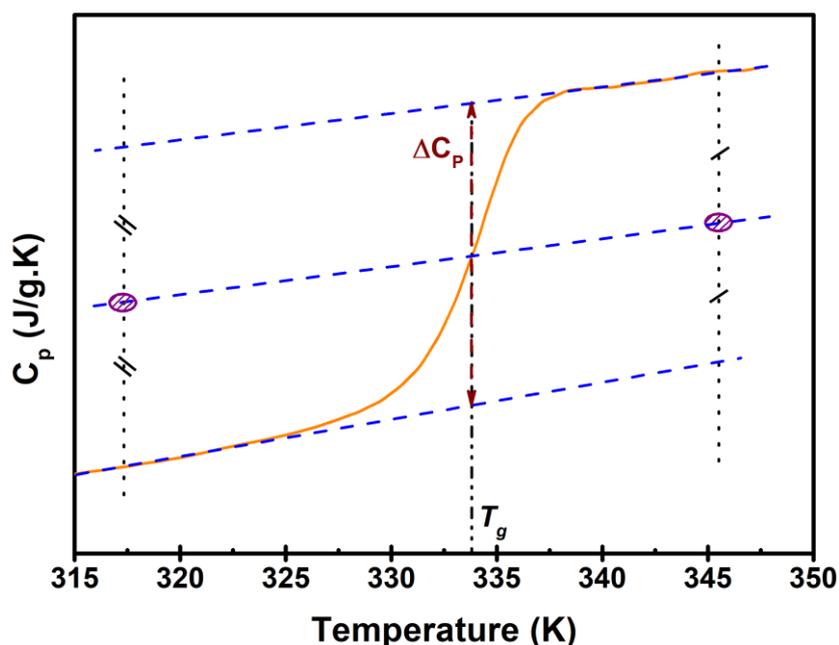
$\Delta t \ll \tau \rightarrow$  glass



**Figure 1. 16.** Schematic illustration of the temperature dependence of the thermodynamic properties of a glass-forming system for three different cooling rates  $\beta_c$ .  $T_{g1}$ ,  $T_{g2}$  and  $T_{g3}$  are the temperatures correspond to the glasses obtained at different cooling rates  $\beta_{c1}$ ,  $\beta_{c2}$ ,  $\beta_{c3}$ , respectively, where  $\beta_{c3} < \beta_{c2} < \beta_{c1}$ .

High cooling rate prevents the possibility of the rearrangement of molecules (cooling rate  $>$  relaxation rate), so the molecules can be frozen rapidly. This fact can explain high speed

cooling rate resulting in a higher  $T_g$ . The glass transition phenomenon can be explained as a discontinuity in the assessment of the enthalpy  $H$  (shown in Fig 1.15). The  $T_g$  will cause a step in the heat capacity plot as a function of temperature and can be estimated by the middle temperature of the occurring variations as shown in Fig 1.17.



**Figure 1. 17.** Temperature dependence of the specific heat capacity of PLLA sample obtained by DSC measurement. The glass transition temperature is determined at the middle point.

The fundamental of glass transition is always a point of debate due to its complexity that includes the equilibrium, thermodynamic and kinetic factors related to the nature of the material. The physical properties of the materials, particularly in the case of amorphous polymers, drastically change at  $T_g$ . Therefore, any physical characteristics (i.e. chain flexibility, stiffness including steric hindrance, polarity, branching) and/or any other impacts (i.e. crystallization, crosslinking) that affect the chain mobility can also influence the value of  $T_g$ .

### 1.8.2 Theoretical approaches of the glass transition

Recently, various theories and models have been developed to explain the fundamental nature of glass transition in the glass-forming systems. Unfortunately, there is no generally accepted theoretical approach that can successfully describe all phenomena of

the glass transition. Nevertheless, theories based on the thermodynamic and kinetic aspects used in this work are briefly introduced.

### 1.8.2.1 Free volume model

The free volume theory proposed by Turnbull and Cohen [82] and Doolittle [97] was based on the following assumptions:

- free volume is continuously redistributed without any expense of local free energy for this redistribution, and
- molecular mobility is associated with movement of atoms or molecules into voids of approximately equal or greater size than that of molecular dimensions.

Motions of the molecules, resulting the redistribution of the free volume  $V_f$ , can be defined as the difference between the total specific volume  $V$  and an occupied volume  $V_0$ , i.e.  $V_f = V - V_0$ . The relation between viscosity and free volume was developed by Doolittle as follows [97]:

$$\ln \eta = \ln A + \frac{B(V - V_f)}{V_f} \quad (1.3)$$

in which  $\eta$  corresponds to viscosity,  $A$  and  $B$  are the fitting parameters,  $V$  is the total specific volume and  $V_f$  is the free volume. The average free volume is  $v_f = V_f / N$ , where  $N$  is the number of molecules. The fundamental assumption from Cohen and Turnbull [83] is based on the linear temperature dependency of average free volume (Eq 1.4). They proposed that redistribution of the free volume in viscous liquids takes place without any charge of the local energy. Thus, molecular motion is associated to the mobility of space which has a sufficient large volume to retain a molecule towards the redistribution of free volume.

$$V_f(T) = \alpha_T \bar{V}_m (T - T_0) \quad (1.4)$$

where  $T_0$  is the temperature at which the average free volume is zero,  $\alpha_T$  is the thermal expansion coefficient and  $\bar{V}_m$  is the mean molecular volume. According to this hypothesis, the free volume model predicts that while the temperature approaches  $T_0$ ,  $v_f$  evanesces, so then the molecular diffusion is stopped. When the contraction occurs, the packing density increases causing a reduced free volume and molecular mobility with the decrease in temperature.

Furthermore, these translational diffusive motions freeze at the glass transition. As a consequence, the free volume approach can be used to clarify strong temperature dependence of relaxation mechanisms like the segmental ( $\alpha$ ) relaxation or viscosity.

### **1.8.2.2 The Adam-Gibbs's model: configurational entropy**

The configurational entropy  $S_c$  was proposed by Adam and Gibbs [85] based on reduction of the relaxation time with decreasing temperature towards the glass transition. The relaxation process of glass-forming liquids takes place by cooperative rearrangements of the molecular groups. Such a motion is only possible if the given number of neighboring molecular groups are dependently in motion. In other words, the rearrangements above energy barriers of molecular units must be “cooperative”, involving a number of repeating units that necessarily increases with decreasing temperature. In addition, the systems can be divided into independent and equivalent regions, so called “cooperative rearranging region” (CRR). CRR is defined as the smallest region that can be exposed a new configuration without any essential simultaneous configurational change. The segmental relaxation is based on the rearrangement of the molecules in the CRRs.

### **1.8.2.3 The concept of cooperativity**

Based on the Adam and Gibbs's assumptions, the system can be divided into  $N$ -independent and equivalent subsystems obtained by  $z$  molecules; also,  $n$  of these regions are assumed to be in the thermodynamic state which leads to cooperative rearrangements. The isothermal-isobaric partition function of the system can be defined as [85]:

$$\Delta(z, P, T) = \sum_{E,V} w(z, E, V) \exp\left(-\frac{E}{k_B T}\right) \exp\left(-\frac{PV}{k_B T}\right) \quad (1.5)$$

in which  $w$  is the degeneracy of energy level  $E$  and volume  $V$  of the subsystems, and  $k_B$  is the Boltzmann constant. If eq 1.5 is limited to the values of  $E$  and  $V$ , the fraction ( $f$ ) of CRRs which involving transitions can then be estimated relatively [84]. The probability of cooperative transition  $W(T)$  with  $z$  molecules, which is proportional to the  $f$  of CRR, can be expressed as:

$$w(T) = A \exp\left(-\frac{z\Delta\mu}{k_B T}\right) \quad (1.6)$$

where  $A$  is a frequency factor which is negligibly dependent on temperature, and  $\Delta\mu$  is the difference between the chemical potential of the molecules in cooperative region and the molecules in the non-cooperative region. While the relaxation time  $\tau$  is related to the transition probability and the relaxation is proportional to the size of CRR region, the temperature dependence of relaxation time in a glass-forming liquid (approaching glass transition) is then expressed as:

$$\tau(T) = \tau_0 \exp\left(\frac{C}{TS_c}\right) \quad (1.7)$$

where  $\tau_0$  and  $C$  are constants,  $S_c$  corresponds to molar configurational entropy of the macroscopic sample. Although this equation depicts the relation between relaxation time and the configurational entropy, the configurational entropy is not experimentally accessible. Therefore,  $S_c(T)$  can be linked to the change in the heat capacity  $\Delta C_p \sim 1/T$  resulting  $S_c(T) \sim (T - T_0) / TT_0$  by using thermodynamic considerations.

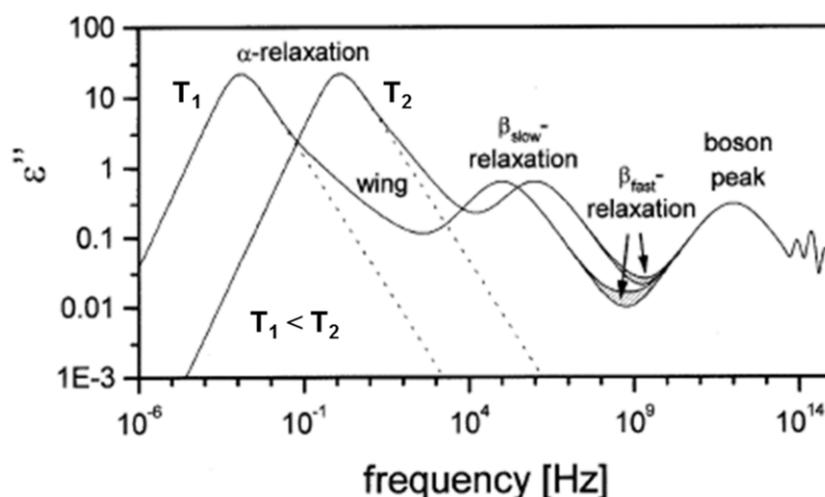
### **1.9 Relaxation phenomena and molecular dynamics**

Different molecular motions exist below and above the glass transition temperature in glass-forming systems. The type of polymer relaxation processes depends on the size of the segments that are related to the molecular movements. These movements can be separated in two main motions:

- ◆ the movement of long chain segments, and
- ◆ the local movement of group of molecules in the chain segment, so called “localized motion”.

Deceleration of the molecular movements in glass-forming liquids broadens the relaxation time or frequency interval, and the time shifts to longer interval at  $T_g$ . The information about the dynamics of these motions can be obtained by measuring the dielectric responses of the sample; i.e. the complex dielectric function ( $\varepsilon^*$ ) or the complex conductivity ( $\sigma^*$ ). The corresponding response of a time dependent dielectric function  $\varepsilon(t)$  to outer alternating electric field  $E(t)$  is called a relaxation (Fig 1.18). The relaxation function is measured as the response to external perturbation based on time or frequency. The

characteristic relaxation time  $\tau$  and/or the relaxation rate  $\nu \sim 1/\tau$  of these motions are measured as a function of temperature by dielectric spectroscopy technique [98]. Therefore, the glass transition temperature is conventionally determined on the basis of the time scale, in which the structural relaxation ( $\alpha$ -relaxation) time  $\tau_\alpha$  reaches 100s.



**Figure 1. 18.** Schematic illustration of dielectric loss spectra in the frequency range  $10^{-6} - 10^{15}$  Hz at temperatures  $T_1$  and  $T_2$  [99]. The dotted lines correspond to the slope at high frequency flank of  $\alpha$ -relaxation peaks.

The characteristic dielectric loss spectra in broad range of frequency at two different temperatures are represented in Fig 1.18, where different relaxation processes in glass-forming systems are depicted. A slow process called  $\alpha$ -relaxation (primary relaxation) is generally observed at low frequency range. It is well known that many glass-forming systems show a  $\beta$ -relaxation and/or an inflection at the high frequency side of  $\alpha$ -relaxation, usually called wing [100, 101]. The secondary process named  $\beta$ -relaxation or  $\beta_{\text{slow}}$  is associated with intramolecular fluctuation and can be observed at higher frequency range. The  $\beta$ -relaxation processes can be attributed to the internal change of the molecular conformation, i.e. a molecular side group motion in a polymer. Some other responses can be observed in the higher frequency range like a peak called boson peak, which corresponds to the vibrational properties of the systems. The  $\alpha$ -relaxation is associated with the segmental movements and is based on the atrophied diffusion of conformational changes as like gauche-trans transition throughout the polymer chain [98]. Such a conformational changes influence the bond length

and the bond angles, thus increases the probability of a neighboring movements that undergoes a cooperative motions. Therefore, the  $\alpha$ -relaxation is a cooperative phenomenon that can be defined as a conformational rearrangement in the amorphous domain which can take place without affecting rearrangements in the surrounding. The cooperative character of the  $\alpha$ -relaxation is described by the size of the cooperative rearranging region (CRR) in the Adam and Gibbs approach [85]. The cooperative molecular motion occurs around the crossover temperature  $T_c$ , and the CRR size increases with decreasing the temperature to the glass transition temperature. Such an increase in the CRR size is linked to the evident characteristics of the relaxation dynamics around the glass transition, i.e. the non-Arrhenius dependence of the  $\alpha$ -relaxation time  $\tau$  [98].

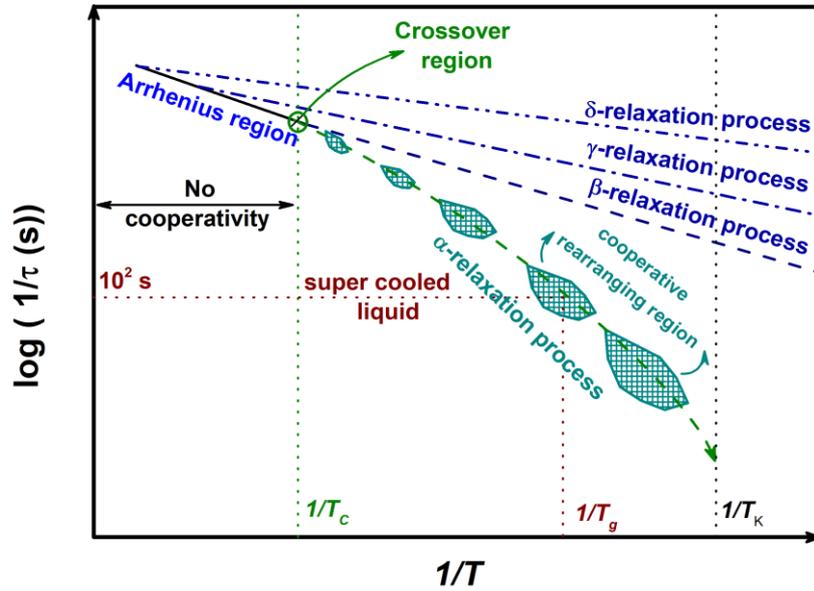
### **1.9.1 The non-Arrhenius temperature dependence**

Although several work proposed the numerous theories, findings and understanding the universal features of relaxation phenomena in glassy systems are challenging [102, 103]. The study of the dynamics of glass-forming liquids is attractive topic to find an explanation of the Arrhenius to non-Arrhenius temperature dependence of the  $\alpha$ -relaxation time  $\tau$  (or viscosity  $\eta$ ) of the glassy polymers while approaching the glass transition temperature. The temperature dependence of the relaxation dynamics can be studied by differential scanning calorimetry (DSC), modulated temperature DSC (MT-DSC), fast scanning calorimetry (FSC), dielectric relaxation spectroscopy (DRS). These techniques allow us to investigate this temperature dependency as schematically demonstrated in Fig 1. 19.

The liquids at very high temperature exhibit Arrhenius temperature dependence, which can be explained by an Arrhenius law:

$$\tau(T) = \tau_{\infty} \exp\left(\frac{E_a}{k_B T}\right) \quad (1.8)$$

where  $\tau_{\infty}$  is the relaxation time at infinite temperature,  $E_a$  is the activation energy required for molecular motions and  $k_B$  is the Boltzmann constant ( $k_B = 1.38 \times 10^{23} \text{ J.K}^{-1}$ ). At higher temperature, molecular motions occur with the coupling of  $\alpha$ - and  $\beta$ - relaxations.



**Figure 1. 19.** Schematic illustration of the temperature dependence of the relaxation ( $\alpha$ ,  $\beta$ ,  $\gamma$ , and  $\delta$ ) processes in a glass-forming liquid during cooling. The cross-over temperature ( $T_c$ ), is the temperature at which the  $\alpha$ - and  $\beta$ - processes combine each other. The Arrhenius-like behavior is depicted at high temperature.  $T_g$  and  $T_k$  are the glass transition and Kauzmann’s temperature, respectively.

However, when the temperature is decreased to the crossover temperature  $T_c$ , these relaxation processes can be separated. The  $\beta$ -relaxation can be still explained by an Arrhenius behavior, the  $\alpha$ -relaxation becomes cooperative motions, and can be described by the Adam and Gibbs theory. The temperature decrease of the  $\alpha$ -relaxation between  $T_c$  and  $T_k$  exhibits the non-Arrhenius behavior (also called super-Arrhenius behavior) which can be explained by the empirical Vogel-Fulcher-Tamman (VFT) equation [104-106] as follows:

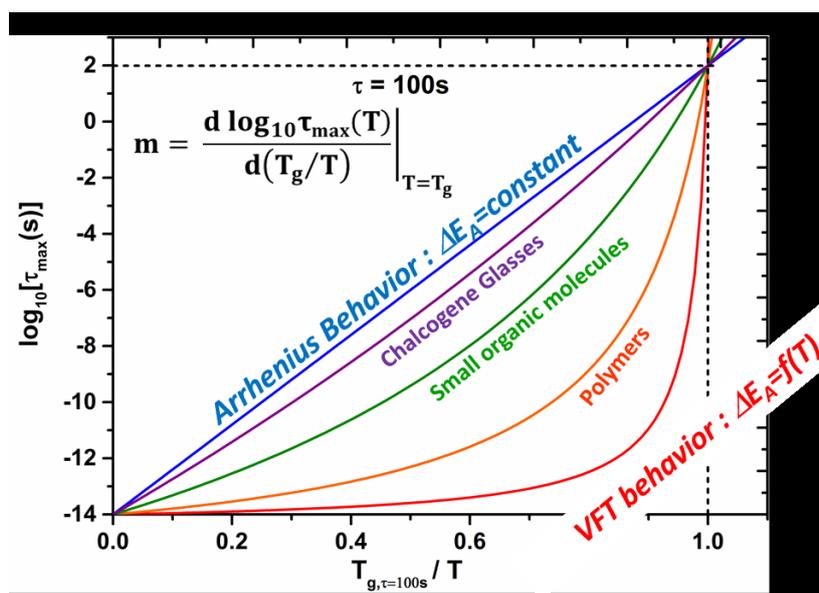
$$\tau(T) = \tau_0 \exp\left(\frac{D_T T_0}{T - T_V}\right) \quad (1.9)$$

where  $\tau_0$  is the relaxation time at infinite temperature,  $D_T$  is the strength parameter related to the fragility, and  $T_V$  is the characteristic temperature below  $T_g$ , so called the Vogel temperature. In addition to the  $\alpha$ -relaxation processes, the CRR size is increased by decreasing temperature from  $T_c$  to  $T_k$ . The VFT law can explain the temperature dependence of the relaxation time of glass-forming liquids up to  $T_c$ . However, Ngai [107] presented that the most of the glass-forming systems, in particular polymer systems, have two differentiated temperature dependence of the relaxation processes at the crossover point. Therefore, two

VFT equations are required for good fitting of relaxation data in the temperature range from  $T_c$  to  $T_g$ . Furthermore, the temperature dependence of the  $\alpha$ -relaxation has been studied by using derivative method that proposed by Stickel et al. [108] confirming the presence of the crossover regions. Stickel et al. defined the association of the VFT law in glass-forming liquids with the linearity of the correlation function  $\Phi_T$  as a function of  $T$ . The Stickel's plot indicates the  $\Phi_T$  versus  $1/T$  whereas  $\Phi_T = [d \log_{10} \tau / d (1/T)]^{-1/2}$ . The non-Arrhenius temperature dependence of the  $\alpha$ -relaxation time has been extensively investigated to classify the glass-forming liquids based on their relaxation time around  $T_g$ . The evaluation from Arrhenius to non-Arrhenius behavior was proposed in the terms of "fragility" to define the degree of the deviation from the Arrhenius dependence [109-111]. Fragility  $m$  is used to estimate the dynamic properties and movements of a glass-forming liquids when temperature approaches the glass transition. It shows the steepness of the temperature variations of viscosity  $\eta$  or relaxation time  $\tau$  approaching  $T_g$ , and is expressed as follows:

$$m = \left[ \frac{d(\log \tau)}{d\left(\frac{T_g}{T}\right)} \right]_{T=T_g} \quad (1.10)$$

where  $m$  is the fragility index,  $T$  is the temperature,  $T_g$  is the glass transition temperature and  $\tau$  is the relaxation time. The value of  $m$  gives information about the categories "strong" or "fragile". The systems which show a strong Arrhenius temperature dependence of  $\tau$  (Arrhenius behavior) are called "strong". Meanwhile, the systems which present a severely non-Arrhenius temperature dependence of  $\tau_\alpha$  (non-Arrhenius or VFT behavior) with steep variations are known as "fragile". The  $m$  value is usually between  $16 \leq m \leq 200$ . The higher value of  $m$  indicates the more fragile systems which show deviation from strong systems, as shown by Angell's plot in Fig 1.20.



**Figure 1. 20.** Angell’s plot presenting the variations of  $\tau$  as a function of the normalized temperature  $T_{g, \tau=100s} / T$  from Arrhenius to non-Arrhenius behavior.

### 1.9.2 The non-exponential behavior

Due to the molecular movements at equilibrium, a simple linear exponential function  $\exp(-t/\tau)$  is not sufficient to explain the isothermal time dependence of the  $\alpha$ -relaxation in liquids. Such behavior can be observed by the time or frequency dependence of the equilibrium liquids measurements, i.e. mechanical [112], dielectric [113] and light scattering [114]. The relaxation or correlation function monitored generally slower decays than the exponential function, so this behavior represents “non-exponential” behavior. The same molecular movements are associated with the structural recovery, therefore the time dependence of the structural relaxation function under non-equilibrium conditions is called also “non-exponential”. After long relaxation time, the systems relax completely and the correlation function decay approaches to zero, this corresponds to the  $\alpha$ -relaxation of the system. Furthermore, due to the decrease in temperature, the  $\alpha$ -relaxation process decelerates and becomes non-exponential near the glass transition. On the other hand, the configurational entropy theory showed that each CRR has unique relaxation time. Therefore, the correlation function in a broad distribution of relaxation time  $\tau$  is well and simply expressed by a stretched exponential function  $\Phi$ , so called Kohlraush-Williams-Watt (KWW) function [115, 116]:

$$\Phi(t) = \exp\left[-\left(\frac{t}{\tau_{KWW}}\right)^{\beta_{KWW}}\right] \quad (1.11)$$

where  $\beta_{KWW}$  is the stretch exponent ( $0 \leq \beta_{KWW} \leq 1$ ) corresponds to the distribution of the relaxation time,  $\tau_{KWW}$  is the relaxation time.  $\beta_{KWW}$  is inversely proportional to the distribution of relaxation time as when  $\beta_{KWW}$  is near to 0, the distribution is very broad. In addition, when  $\beta_{KWW}$  is close to 1, the time distribution can be characterized by a simple exponential so called Debye relaxation that is observed in liquids above the melting point. Debye relaxation was firstly reported in 1863 to explain the mechanical creep in glassy fibers [115] and was then studied by Williams and Watts in 1970 to investigate dielectric relaxation in polymer systems [116]. The KWW law can be used for different physical and mathematical calculations of models based on distribution of relaxation times to complex correlated processes.

### 1.10 Dynamic heterogeneity and cooperative movements

There are some important concepts that have been universally accepted and in which the experimental results are linked to the investigation of vitrification and its dynamics. The most well-known studies are the notions of cooperativity and the spatially heterogeneous dynamics, named “dynamic heterogeneity”. The concept of the dynamic heterogeneity is applied to find an answer for why there are severe variations of experimental parameters like relaxation time and viscosity of glass-forming liquids towards glass transition [117-119]. Ediger [118] proposed that the dynamic heterogeneity may be expressed by the different regions (few nanometers away from each other) in a liquid including sub-ensembles of particles that show momentarily increased or diminished mobility relative to the average mobility. If these particles have higher mobility than the average, they are defined as “mobile” or “fast” particles. Those particles with lower mobility are considered to be “immobile” or “slow” particles. Dynamic heterogeneity in the glass-forming liquids is not static as it improves over time. The notion of dynamic heterogeneity exudes the dynamically correlated movements of the particles in the regions. Therefore, molecular cooperativity associated with the glass transition is intimately related to the dynamic heterogeneity [120-124]. Indirect methods, i.e. multipoint dynamic susceptibility function [125-127] and thermodynamic methods based on the fluctuation-dissipation theory [123, 128], are currently used for understanding the dynamic heterogeneity. These approaches lead to estimate the length scale at  $T_g$  that is

approximately from 1 nm to 4 nm for different glass-forming liquids [129]. It has been found that the dynamics of the glass-forming liquids is dynamically heterogeneous towards the calorimetric glass transition temperature  $T_g$ . The discovery of the dynamical heterogeneity well defines the deviation from the exponential behavior of the glass-forming systems responses. Furthermore, the variations in the temperature dependence of translational or rotational diffusion in the systems approaching to the glass transition can be also defined by the dynamic heterogeneity existence. Thus, the approaches have been focused on the notion of the heterogeneous dynamics of the  $\alpha$ -relaxation processes in the liquid-like state for better understanding the nature of the molecular mechanisms associated to the glass transition. Therefore, the majority of the research groups focus on the length scale of dynamic heterogeneity and the cooperativity as a function of time and temperature for polymer glass-forming liquids. Two independent models are mostly used to estimate the length scale. One model was proposed by Berthier et al. [125] on the basis of the higher order correlation function, the other one was reported by Donth [130] based on the fluctuation dissipation theory.

#### **1.10.1 Length scale from four-point dynamic susceptibility function**

Since the  $\alpha$ -relaxation exhibits both super-Arrhenius and non-exponential behavior towards the glass transition temperature, the existence of the length scale associated to the coexistence of molecules with essentially different relaxation time is still widely investigated. Dynamical heterogeneity is spatially organized into fast and slow regions with a typical size. To define the quantity in dynamics, the two-point correlation function is used [125] and is generally expressed by the KWW (a stretched exponential) function to define the  $\alpha$ -relaxation in glass-forming systems. However, the two-correlation function cannot individuate the homogenous and heterogeneous movements in one region of a glass-forming liquids. Therefore, higher order correlation function “four-point dynamic susceptibility” of the density fluctuation is used to investigate the behavior and magnitude of the dynamically correlated molecules in glass-forming liquids (i.e. repeating units in the case of polymers). Four-point dynamic susceptibility corresponds to the spatial correlation of localized particles, as the evident location is associated to the correlated motion of the particles occupied by neighboring particles [131, 132]. When two-point correlation function involves the variables

at two times for the dynamics description, the four-point correlation function expresses the spatial and temporal fluctuation of dynamic heterogeneity [120]. Therefore, the four-point susceptibility  $\chi_4(t)$  is estimated by the integration of the probability of a similar event that occurs at a distance  $r$  at the same time interval  $t$  ( $C_4(r, t)$ ) as [133]:

$$\chi_4(t) = \int C_4(r, t) dr \quad (1.12)$$

where  $\chi_4(t)$  corresponds to the maximum value at the relaxation time  $\tau_\alpha$  of glass-forming liquids. Interactions between regimes play a role on a longer time scale, then these regimes are separated in early and late regimes due to the elasticity theory. Later then, because of the approach and departure from the plateau of the four-point susceptibility function, the relaxation time  $\tau_\alpha$  is reached, and  $\chi_4(t)$  presents a peak with a maximum, which is proportional to the volume where correlation movements occur. The number of correlated molecules  $N_{COOR,4}$  in an average relaxation time  $\tau_\alpha$  can be estimated as [120, 122, 125]:

$$N_{COOR,4} = \max_t \{X_4(t)\} \quad (1.13)$$

Due to the impossibility of an experimental measurements of  $\chi_4(t)$ , Berthier et al. [125] proposed a new model to estimate a typical length scale of the dynamic heterogeneity on the basis of the temperature dependence. They modified the four-point susceptibility function to the three-point dynamic susceptibility  $\chi_T(t)$  which is experimentally accessible [125]. The  $\chi_T(t)$  exhibits a peak at  $t \sim \tau_\alpha$  from which the number of molecules  $N_C$ , whose dynamics on the time scale of the  $\alpha$ -relaxation are correlated, can be estimated [134]:

$$N_C = \frac{N_A}{m_0} \frac{k_B}{\Delta C_P} T^2 \max(\chi_T(t))^2, \text{ or } N_C = \frac{N_A}{m_0} \frac{k_B}{\Delta C_P} T^2 \max\left(\frac{\delta\varepsilon}{\delta T}\right)^2 \quad (1.14)$$

where  $N_C$  is the number of the dynamically correlated segments (repeating units in polymers),  $m_0$  is the molecular weight of the repeating units,  $N_A$  is the Avogadro's number,  $\Delta C_P$  is the variation in heat capacity between liquid and glassy states,  $k_B$  is the Boltzmann constant, and  $\varepsilon$  is the dielectric relaxation function. When the dielectric relaxation function shows a Kohlrausch form with a stretch exponent, the number of dynamically correlated units can be estimated by a simpler form or by eq. 1.14 as Capaccioli et al. proposed [135]:

$$N_C = \frac{N_A}{m_0} \frac{k_B}{\Delta C_p} \left( \frac{\beta_K}{e} \right)^2 \left( \frac{d \ln \tau_\alpha}{d \ln T} \right)^2 \quad (1.15)$$

where  $\Delta C_p$  is the variation in heat capacity between liquid and glassy states,  $m_0$  is the molecular weight of the repeating units,  $N_A$  is the Avogadro's number,  $k_B$  is the Boltzmann constant,  $\beta_K$  is the stretch parameter,  $e$  is the Euler's number and  $\tau_\alpha$  is the relaxation time of  $\alpha$ -process. As consequence,  $N_C$  increases with decreasing temperature and increase of  $\tau_\alpha$  as a function of temperature  $T$  and relaxation time  $\tau_\alpha$ , respectively.

In this work, the number of dynamically correlated systems is calculated using eq. 1.15. Different parameters in eq. 1.15 can be usually exported from the results of two experimental techniques that are DRS and MT-DSC.

### **1.10.2 Length scale from temperature fluctuation: Donth's approach**

Donth proposed a simple method based on the temperature fluctuation in subsystems (CRR) in order to calculate the length scale at the calorimetric glass transition [129, 136]. As explained in the Adam-Gibbs's theory, the CRRs are definable independent subsystems that demonstrate thermodynamic fluctuations [85]. Therefore, CRR represents an independent relaxation from the rest of the system over its own relaxation time  $\tau$  as well as own glass transition  $T_\alpha$ . In addition, the dispersion in relaxation time can be linked to the distribution of  $T_\alpha$  between these subsystems. Therefore, an average dynamic glass transition  $\langle T_\alpha \rangle$  of all subsystems can be defined as the conventional calorimetric glass transition temperature  $T_g$ . The density  $\rho$ , the temperature  $T$ , the entropy  $S$  and the energy  $E$  of the subsystems can be different in each equal sub-volume of the glass-forming liquids. The thermodynamic fluctuations were limited to slow degree freedom of the  $\alpha$ -relaxation and the width of the relaxation times of  $\alpha$ -processes [137]. Therefore, a characteristic length of glass transition can be estimated from experimental data.

Donth assumed that each CRR has partial structural entities with own frequency and local free volume that fluctuate during time as frequency is the function of the partial free volume. CRR is divided into the equivalent partial systems which are considered as statically independent but cooperatively associated by free volume redistribution. Therefore, Donth considered that the partial functions for each entities are independent and cooperative

movements are caused by the distribution of the partial free volume. Moreover, these partial subsystems can be investigated by its frequency and free volume and involved populations that are linked to a temperature. Then, an average temperature fluctuation  $\delta T$  confirmed in a CRR between the different populations.

The independent and equivalent free volume is thermodynamically in the metastable equilibrium where the fluctuation in variables can be defined by Gaussian fitting. The energy fluctuation is dependent on the isochoric heat capacity ( $J.K^{-1}$ ), the freedom degree of the  $\alpha$ -relaxation processes and vibrational motion associated with glass below  $T_g$  [138]. Therefore, the relation between the average  $T_g$  and the thermal heat capacity can be defined as:

$$(\delta T_g)^2 = k_B T_g^2 \frac{\Delta C_V}{C'_{V,\alpha}} \quad (1.16)$$

and/or

$$\frac{\Delta C_V}{C'_{V,\alpha}} \approx \Delta \left( \frac{1}{C_V} \right) = \frac{1}{C_{V,glass}} - \frac{1}{C_{V,liquid}} \quad (1.17)$$

By assuming that the variation of  $1/C_P$  is identical with that of  $1/C_V$ , then:

$$(\delta T_g)^2 = k_B T_g^2 \Delta \left( \frac{1}{C_P} \right) \quad (1.18)$$

can be expressed. To calculate a length scale of cooperativity, the heat capacity  $C'_{P,\alpha}$  ( $J.K^{-1}$ ) can be modified in terms of the specific heat capacity in unit of  $J g^{-1} K^{-1}$  as follows:

$$C'_{P,\alpha} = C_{P,\alpha} \rho V_\alpha \quad (1.19)$$

in which  $C_{P,\alpha} = C_{P,T>T_g} - C_{P,T<T_g}$  is the change in the specific heat capacity step at the glass transition. Thus, the characteristic length scale of the glass transition  $\xi_\alpha$  and the volume can be expressed as:

$$\xi_\alpha = \sqrt[3]{V_\alpha} = \sqrt[3]{\frac{\Delta C_P^{-1} k_B T_g^2}{\rho (\delta T)^2}} \quad (1.20)$$

where  $\Delta C_p^{-1}$  is the variation in the inverse of the isobaric heat capacity for liquid and glass at the dynamic glass transition temperature  $T_\alpha$ ,  $k_B$  is the Boltzmann constant,  $\rho$  is the density at  $T_\alpha$  and  $\delta T$  is the temperature fluctuation in a CRR at  $T_\alpha$ .

Correspondingly, the number of structural units in a CRR ( $N_\alpha$ ) can be estimated as:

$$N_\alpha = \frac{N_A k_B T_\alpha^2 \Delta C_p^{-1}}{m_0 (\delta T)^2} \quad (1.21)$$

where  $N_A$  is the Avogadro's number and  $m_0$  is the molar mass of repeating unit.

The parameters of the Donth's approach,  $T_\alpha$ ,  $\delta T$  and  $\Delta C_p^{-1}$  can be exported from calorimetric techniques such as DSC and MT-DSC. The cooperativity length  $\xi_\alpha$  at the glass transition estimated by Donth's approach for glassy polymers is usually in the range of 1-3.6 nm [139-142].

### 1.10.3 The cooperativity size in a wide temperature and frequency range

The classical Donth's model proposed the CRR size estimation in only narrow temperature and frequency range close to the calorimetric  $T_g$ . However, the combination of two experimental techniques (MT-DSC and DRS) allow us to estimate the cooperativity length in a broad temperature and frequency range [142, 143]. There are several common and complementary notions of the glass transition used in these two techniques:

- ◆ the dynamic glass transition can be monitored by applying a periodic perturbation to the sample, i.e. a modulated temperature ramp in MT-DSC and a periodic electric field in DRS. Therefore, such techniques allow us to measure the complex quantities of the heat capacity  $C_p^*$  by MT-DSC and the dielectric permittivity  $\varepsilon^*$  by DRS involving real ( $C_p'$  and  $\varepsilon'$ ) and imaginary ( $C_p''$  and  $\varepsilon''$ ) parts, respectively;
- ◆ due to the linear response theory, both real and imaginary quantities can be directly compared;
- ◆ in case of MT-DSC, the dynamic glass transition can be estimated by a heat capacity step  $\Delta C_p$  in the real part, while the relaxation peak can be observed in the imaginary part;

- ♦ in case of DRS, the dynamic glass transition can be determined from the dielectric strength  $\Delta\varepsilon$ , and a peak of  $\varepsilon''$  in the imaginary part.

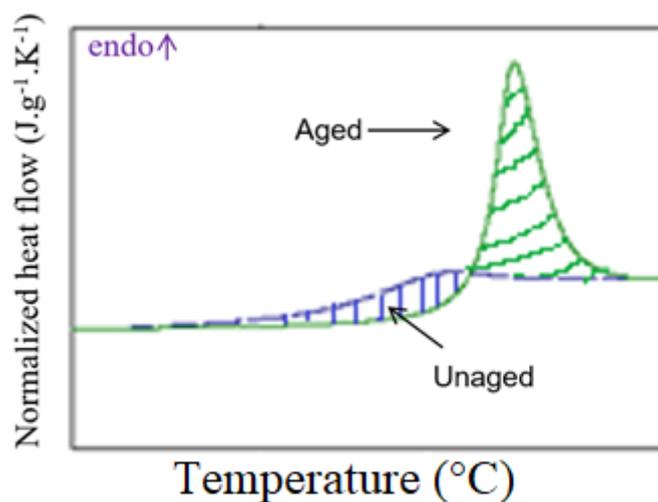
Therefore, both  $\Delta C_p$  and  $\Delta\varepsilon$  correspond to the amorphous phase species at the dynamic glass transition temperature. So, the glass forming liquids exhibit a relaxation time distribution function at each temperature step. As the consequence, the temperature fluctuation  $\delta T$  is associated to the standard deviation  $\sigma_T$  of the Gaussian peak function, so the average temperature fluctuation  $\delta T$  can be obtained from the imaginary part  $C_p''(T)$  of complex heat capacity as well as from the imaginary part  $\varepsilon''(T)$  of the complex dielectric permittivity. In DRS analysis, the dynamic glass transition  $T_\alpha$  is determined from the maximum of the peak in DRS spectra. The parameters of the Donth's approach,  $T_\alpha$ ,  $\delta T$  and  $\Delta C_p^{-1}$  can be obtained by these two techniques and combined to extend Donth's approach to calculate the temperature dependence of the CRR size as well as the cooperativity degree  $N_\alpha$  in a wide temperature range (from Eq 1.20 and 1.21, respectively).

### **1.11 Structural relaxation of glassy polymers**

Besides the Arrhenius and non-Arrhenius temperature dependences of the  $\alpha$ -relaxation process in glass-forming liquids, the structural relaxation takes place when the temperature approaches the glass transition temperature and below.

#### **1.11.1 Physical aging concept**

Even though the glassy form seems to be "frozen" on the measurement scale below glass transition temperature, a molecular mobility still exists [92, 93]. Therefore, when the glass-forming liquids are held below their respective glass transition temperature, so called physical aging (the structural relaxation) will occur. Such a structural relaxation is the convergent evolution of the thermodynamic properties towards equilibrium [93]. Such thermodynamic properties towards equilibrium are associated with a free volume hole diffusion towards surfaces which could be the external ones of the system or the internal ones in the low density region obtained by controlled cooling rate [144-146]. Thus, investigation of structural relaxation in the terms of physical aging gives information about molecular motions/dynamics.



**Figure 1. 21.** Illustration of the heat flows of a glass-forming liquids during physical aging. Green area corresponds to superimposed peak at  $T_g$  (aged sample), blue area corresponds to unaged sample.

The physical aging can be studied by calorimetric investigations. When glass is held isothermally below the glass transition temperature, the following heating curve will exhibit a peak superimposed to the glass transition as shown in Fig 1.21. Such a peak is related to the enthalpy recovery of a glass forming liquids and gives an information about the excess of the energy released during physical aging. Enthalpy recovery of a glass that aged at a temperature  $T_{ag}$  for an aging time  $t_{ag}$  is calculated by the integration of the variations between the measure of aged and rejuvenated sample by following equation [147]:

$$\Delta H (T_{ag}, t_{ag}) = \int_{T_1}^{T_2} [C_P^a(T) - C_P^r(T)]. dT \quad (1.22)$$

where  $C_P^a(T)$  and  $C_P^r(T)$  are the heat capacity of the aged and rejuvenated (unaged) sample, and  $T_1$  and  $T_2$  are the optional temperatures below and above the glass transition temperature, respectively. It is considered that equilibrium is reached for an infinite aging time. Therefore, the expected total enthalpy loss  $\Delta H_\infty$  can be extrapolated from the equilibrium depending on the aging temperature  $T_{ag}$  and the glass transition temperature  $T_g$ , as well as the heat capacity step  $\Delta C_P$  of the sample, as expresses by follows:

$$\Delta H_\infty = \Delta C_P (T_g - T_{ag}) \quad (1.23)$$

### 1.11.2 The fictive temperature concept

Due to the kinetics related with the glass transition temperature and devitrification upon heating,  $T_g$  can be measured by the concept of the fictive temperature  $T_f$ . The fictive temperature is thermodynamically defined as the crossover point of the enthalpies of the glassy and the liquid-like state at non-equilibrium process. This concept has been reported by Tool et Eicitlin in 1931 [148]. The kinetics of the structural relaxation can be also investigated by measuring the fictive temperature as a function of aging time. The fictive temperature concept is also used to examine the cooling rate dependence of the glass transition in order to understand the nature of glass-forming liquids. The kinetic impact of the cooling rate can be observed upon subsequent heating from the glassy state to liquid-like state. When a glass is obtained at  $\beta_{c1}$  and then it is heated at the same rate ( $|\beta_{c1}| = \beta_{h1}$ ), the enthalpy recovery will occur on the same line at the liquid-like state. However, if a glass is formed at  $\beta_{c2}$  ( $\beta_{c2} \ll \beta_{c1}$ ), the glass transition temperature will decrease. Then, a glass is heated at  $\beta_{h1}$  ( $\beta_{h1} \gg \beta_c$ ), the enthalpy recovery exhibits a peak superimposed on the glass transition upon heating. The fictive temperature  $T_f$  can be measured from calorimetric measurements, i.e. DSC and FSC by the Moynihan's equation [149]:

$$\int_{T_1}^{T_f} (C_{P,liquid} - C_{P,glass}) \cdot dT = \int_{T_1}^{T_2} (C_P - C_{P,glass}) \cdot dT \quad (1.24)$$

where  $T_1$  and  $T_2$  are the optional temperatures above and below the glass transition, respectively;  $C_P$  is the heat capacity of the sample at constant pressure,  $C_{P,liquid}$  and  $C_{P,glass}$  are the extrapolated liquid and glassy heat capacity lines of the sample, respectively.

Background history of PLA-based materials and related theoretical approaches have been presented in this chapter. Existing literature focused its attention to the crystalline structure of sc PLA with very little emphasis on the amorphous phase. With this in mind, we established our objectives to understand the structural-property relationship of sc PLA. In Chapter 2 detailed experimental set-ups are presented in order to achieve the mentioned objectives. The effect of two different elaboration techniques on the sc PLA formation was first investigated (Chapter 3). Following this, we investigated the impact of tacticity/chirality on the amorphous species of sc PLA (Chapter 4). And lastly, the glass transition property in terms

of free volume concept (molecular mobility) was linked to the transport property of sc PLA as compared to that of homopolymers and commercial PLA (Chapter 5).

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**CHAPTER 2. EXPERIMENTAL PART**

This chapter presents the detailed description of experimental techniques (calorimetric investigations, microstructure analysis and permeametric analysis) used in this thesis in order to analyze the elaborated samples. Experimental set-ups and protocols applied during the analysis were briefly described.



## 2.1 Materials

PLLA and PDLA were chosen due to the presence of chiral carbon in the skeletal chain of PLA that yields to two stereoregular isomers, PLLA and PDLA. The mixture of these homopolymers by the stereocomplexation reaction allows obtaining a new structure, so called sc PLA. In this work, homopolymers of poly(L-lactide) (PLLA) and poly(D-lactide) (PDLA), and stereocomplex poly(lactide) (sc PLA) were studied (Fig 2.1).

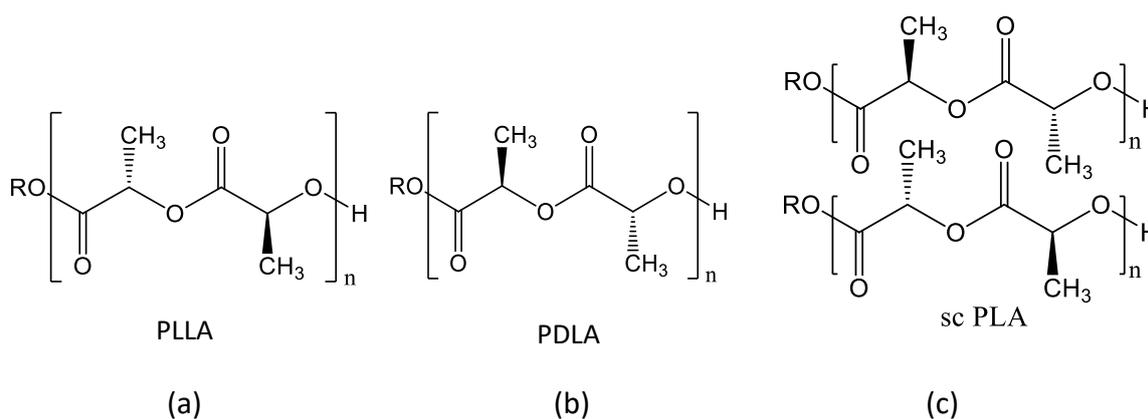


Figure 2. 1. Repeating units of (a) PLLA, (b) PDLA, and (c) sc PLA.

### 2.1.1 Homopolymers

PLLA pellets (NaturePlast Injection grade, France) have the density of 1.25 g/cm<sup>3</sup>. The content of the *D*- isomers is negligible. The melting temperature  $T_m$  is between 170 °C and 180 °C. The number-average and weight-average molecular weight measured by Gel Permeation Chromatography (GPC) are  $M_n = 53$  kDa and  $M_w = 97$  kDa, respectively.

PDLA pellets (NaturePlast PDLA 002, France) have the density of 1.25 g/cm<sup>3</sup> with the content of the *L*-isomer less than 0.5%. The melting temperature  $T_m$  of PDLA is about 175 °C. The number-average and weight-average molecular weight measured by GPC are  $M_n = 56$  kDa and  $M_w = 99$  kDa, respectively. The molar mass of repeating unit (Fig 2.1 (a) and (b)) is  $m_0 = 72$  g/mol. Both PLLA and PDLA homopolymers were dried at  $T = 55$  °C during 3 days before using.

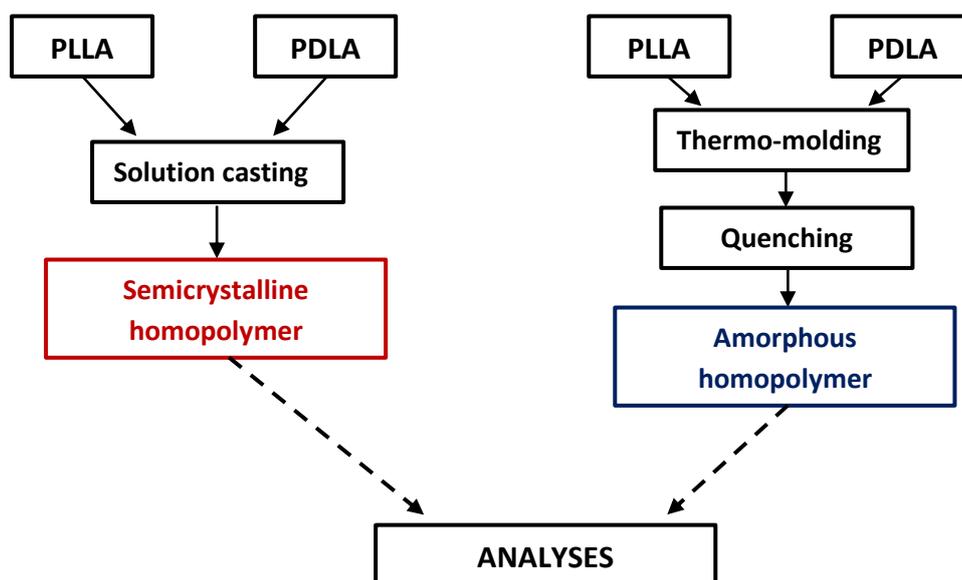
#### 2.1.1.1 Preparation of amorphous homopolymers

After a study of the thermal stability by thermogravimetric analysis (TGA) (the degradation temperature  $T_d = 270$ -400 °C), the dried PLLA and PDLA pellets were thermo-

molded at 1 bar and 180 °C between hot plates during 5 min by using a manual hydraulic press ATLAS™ SPECAC. The temperature was selected above the melting temperature  $T_m$  (ranging from 160 °C to 175 °C [1, 2]) and below the equilibrium melting temperature  $T_m^0$  of PLAs (ranging from 199 °C to 207 °C [3]). The films samples were then rapidly quenched in liquid nitrogen and then dried at 60 °C for 24 h. The thickness of the obtained PLLA and PDLA films was 150 - 250  $\mu\text{m}$ .

### 2.1.1.2 Preparation of semicrystalline homopolymers

Semi-crystalline PLLA and PDLA films were obtained by solution casting. Chloroform (99.8% purity) was used as a solvent. Dried pellets were dissolved in chloroform with the concentration of 3 g/dL at  $T_{amb} = 24 \pm 1$  °C under vigorous stirring during 1h. The prepared solution was poured onto glass Petri dish to obtain free standing films. The solvent evaporation was carried under the fume-hood at ambient temperature ( $T = 24 \pm 1$  °C) for 48h, and the samples were further dried in oven at 120 °C for 12h to remove the solvent completely. The overall preparation process for amorphous and semicrystalline homopolymers is summarized in Fig 2.2.



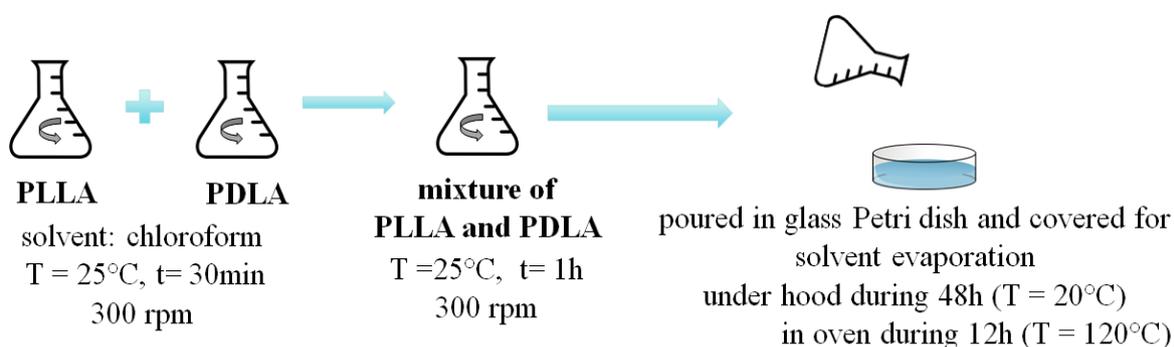
**Figure 2. 2.** Schematic illustration of homopolymer film preparation.

### 2.1.2 Stereocomplex poly(lactide) (sc PLA)

In order to study the influence of elaboration method on the film properties, stereocomplex poly(lactide) (sc PLA) was obtained by two elaboration methods, i.e. solution casting and extrusion process.

#### 2.1.2.1 Solution casting

Sc PLA (also referred as PLLA/PDLA films in some Figures) was prepared by solution casting using chloroform (99.8% purity) as a solvent. In our work, only the influence of solution concentration and PLLA:PDLA ratio was investigated. In order to study the influence of homopolymer concentration on the stereocomplex formation, the solutions with different polymer concentrations (namely 1, 2 and 3 g/dL) were prepared. In this case, the ratio of PLLA:PDLA was maintained constant at 50:50 (w/w). In order to study the impact of homopolymers ratio on stereocomplexation, the solutions with different ratios of PLLA:PDLA were prepared: 50:50, 60:40, 70:30, 75:25, 90:10 and 95:05 ratio of PLLA:PDLA for the concentration of 3 g/dL (this concentration was chosen after careful analysis of the XRD patterns that will be discussed later in Chapter 3). Dried PLLA and PDLA pellets were dissolved in chloroform separately at  $24 \pm 1$  °C. Then, PLLA and PDLA solutions were mixed together under vigorous stirring for additional 1h. The prepared solution was poured onto glass Petri dish (as shown in Fig 2.3). Solvent evaporation was carried out in the fume-hood ( $T \approx 20$  °C) for 48h and then additional drying took place in convection oven ( $T = 120$  °C) for 12h. The obtained free-standing films had the thickness of  $\sim 150$   $\mu\text{m}$ .



**Figure 2. 3.** Schematic illustration of the solvent casting process.

### 2.1.2.2 Quenching

PLLA/PDLA films elaborated by solution casting were thermo-molded at 230 °C and 1 bar between two hot plates during 5 min by using a manual hydraulic press ATLAS™ SPECAC. The temperature was selected close to the equilibrium melting temperature  $T_m^0$  of 235 °C for sc PLA crystals [4]. The samples were then quickly quenched in liquid nitrogen to obtain amorphous films. The thickness of the obtained films was remained  $150 \pm 20 \mu\text{m}$ .

### 2.1.2.3 Extrusion

Blend of sc PLA was prepared by twin screw mixing. PLLA and PDLA pellets were mixed with the ratio of 50:50 (w/w) for extrusion process. The influence of extrusion temperature ( $T_{ext}$ ) and mixing time ( $t_{ext}$ ) on the stereocomplexation of PLA was investigated. In order to investigate the temperature dependence of extrusion, sc PLA was extruded at  $T_{ext} = 180, 190, 200, 210,$  and  $220 \text{ °C}$  during  $t_{ext} = 30 \text{ min}$  with a rotation of 40 rpm. The impact of time was observed by comparing the extruded samples at different time ( $t_{ext} = 15 \text{ min}$  and  $30 \text{ min}$ ) at a fixed temperature ( $T_{ext} = 220 \text{ °C}$ ). The homogeneity of mixing was taken into account by the feeding amount of mixer because of its capacity. The amount of feeding was fixed as 6 g of polymers ( $m_{\text{PLLA}} = m_{\text{PDLA}} = 3 \text{ g}$ ). Fig 2.4 illustrates the overall process for the sc PLA elaboration.

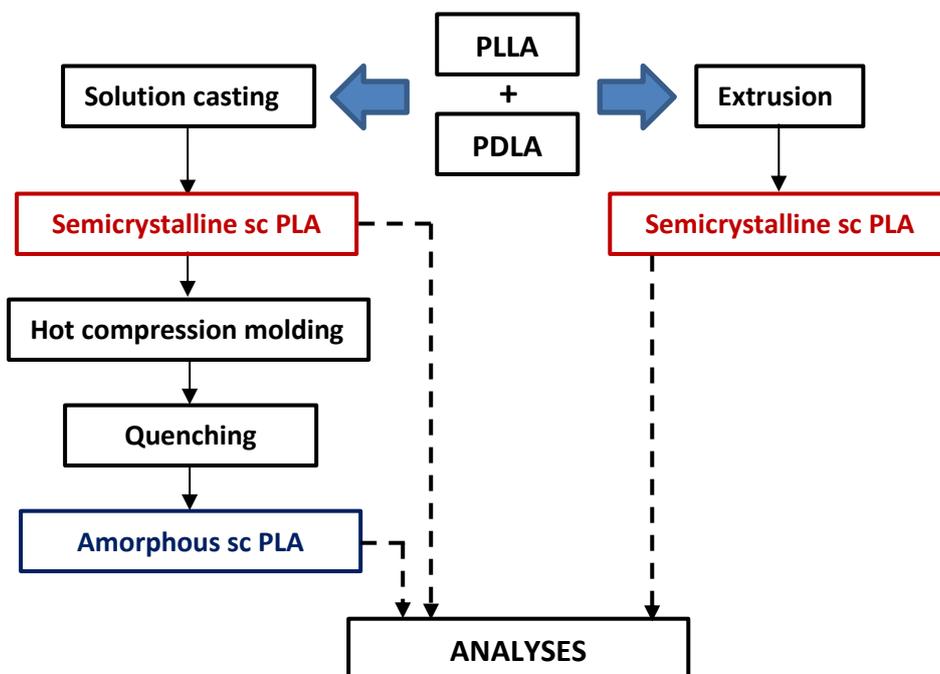


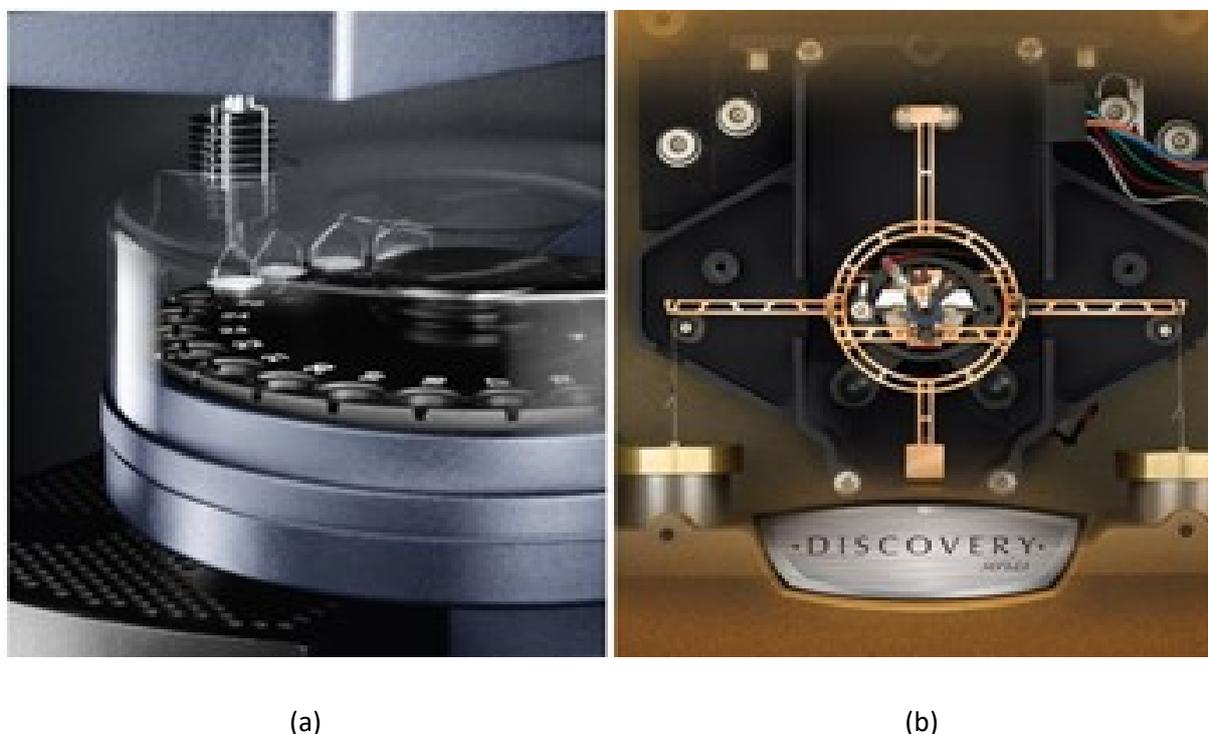
Figure 2. 4. Schematic illustration of the PLLA/PDLA sample preparation.

## 2.2 Instrumentation and methods

The influence of stereocomplexation on the microstructure and the material properties was studied by the microstructure and morphology analysis. In addition, calorimetric investigations were performed in order to understand the fundamentals of short- and/or long-range molecular mobility.

### 2.2.1 Thermogravimetric analysis (TGA)

TGA is a kind of thermal analysis in which the sample weight loss is measured either as a function of increasing temperature, or isothermally as a function of time, under nitrogen, helium, air, other gas, or vacuum. It provides information about different chemical phenomena such as thermal decomposition and physical phenomena including phase transitions, absorption and desorption [5]. It is a beneficial technique for the study of polymer materials, including thermoplastics, thermosets, elastomers, composites, fibers, and coating [6-8]. In this work, the measurements were performed by TGA Discovery (TA Instrument) (shown in Fig. 2.5) to determine the degradation temperature of the samples.



**Figure 2. 5.** Images of (a) sample pan place and (b) balance in the furnace of TGA Discovery.

The samples were placed in the pan (Fig.2.5. a) with the mass around 5 mg and measurements were carried out from 30 °C to 600 °C with a heating rate of 10 °C/min under nitrogen. The degradation temperature ( $T_{deg}$ ) was recorded at the 1% mass loss of the sample.

### 2.2.2 Differential scanning calorimetry (DSC)

DSC is the most used thermal technique that allows us to make qualitative and quantitative measurements. DSC determines the heat flow and the temperature related to calorimetric transitions as a function of temperature and time. DSC provides information about physical and chemical changes including endothermic (sample absorbs energy) and exothermic (sample releases energy) processes of materials during physical transitions which are caused by phase changes, cold crystallization, melting, crystallization, oxidation, and other heat-related changes [9]. Two kinds of DSC exist, so called heat-flux DSC and power-compensated DSC. In this thesis, the calorimetric investigations were performed with a heat-flux calorimeter from Thermal Analysis Instruments (DSC Q100 TA Instruments). The fundamental principle of this DSC lies in the difference between the amount of heat required to increase the temperature of a sample and that of a reference. The measurements of the heat-flux DSC  $T_{zero}^{\text{TM}}$  technology is based on the difference of heat flow between sample, reference, and furnace. In this type of DSC, the sample pan and reference pan are placed in a unique furnace (Fig.2.6).

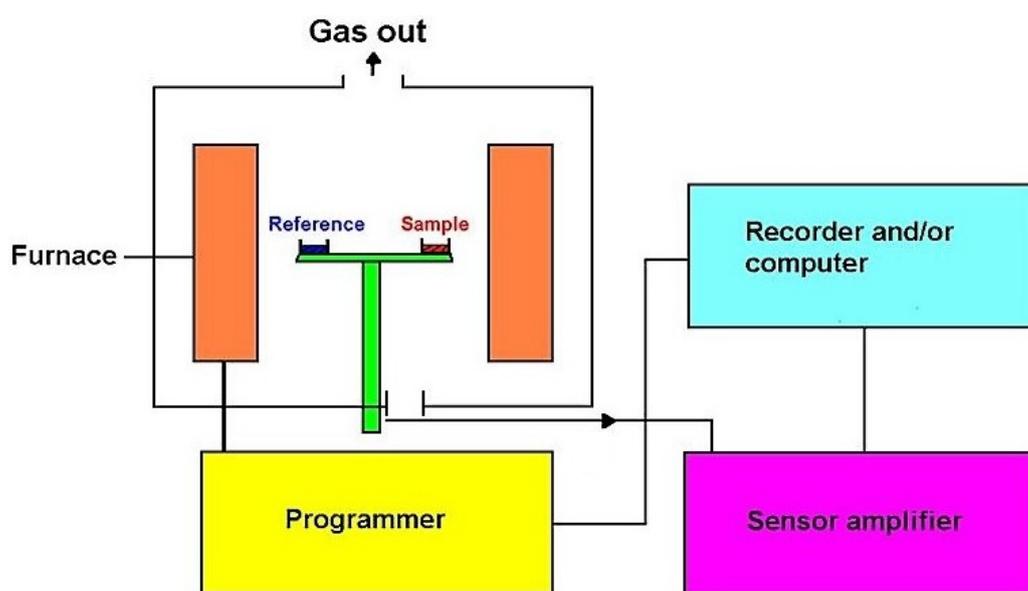


Figure 2. 6. Schematic representation of heat-flux DSC.

The furnace is subjected to the  $T$  changes and heat is transferred to the sample and reference. The temperature is measured continuously, and a differential technique is utilized to detect the heat flow into the sample and to equalize random heat gains and losses between reference and sample. DSC does not allow a direct determination of the heat content. In conventional DSC, the applied temperature ramp is linear:

$$T = T_i + \beta t \quad (2.1)$$

where  $T$  is the temperature at the time  $t$ ,  $T_i$  is the initial temperature and  $\beta$  is the scanning rate. The resulting heat flow  $\Phi$  in the heat transfer  $Q$  is determined as follows:

$$\Phi = \frac{dQ}{dt} \quad (2.2)$$

At the beginning of the measurement, the two thermocouples, i.e. sample and reference, are at the same temperature  $T_0 = T_s = T_r$ , where  $T_0$  is temperature of the location of measurements,  $T_s$  is temperature of sample pan, and  $T_r$  is temperature of the reference pan. During heating, the temperature variation occurs between the sample pan and the reference pan due to the additional weight of sample. Therefore, the variation of heat transfer onto sample ( $Q_s$ ) and reference ( $Q_r$ ) is observed, which is related to the endothermic or exothermic events of the sample. The relation between heat transfer and the variation of temperature can be expressed by a thermal equivalent of the Ohm's law [10]:

$$\Phi = Q_s - Q_r = \frac{\Delta T}{R} \quad (2.3)$$

where  $R$  is the thermal resistance. This variation in temperature is associated to the specific heat capacity of the sample [11]:

$$\Phi = \frac{dQ}{dt} = C \frac{dT}{dt} = c \cdot m \cdot \beta \quad (2.4)$$

where  $C$  is the heat capacity,  $c = C/m$  is the specific heat capacity,  $m$  is the sample mass, and  $\beta$  is the scan rate (heating or cooling rate).

Since, DSC Q100 is based on the  $T_{zero}^{\text{TM}}$  technology and as such technology uses more complex equations of the heat flow, the relations between thermal resistance  $R_{SCP}$  and  $R_{RCP}$ ,

and heat capacities  $C_{SCP}$  and  $C_{RCP}$  of the sample and reference cell platforms, respectively, are taken into account in the heat flow equation. Therefore, the temperatures  $T_s$  and  $T_r$  monitored as a function of a location temperature  $T_0$  can be expressed as follows [12]:

$$\Phi = -\frac{T_s - T_r}{R} + (T_0 - T_s) \left( \frac{1}{R_{SCP}} - \frac{1}{R_{RCP}} \right) + (C_{RCP} - C_{SCP}) \frac{dT_s}{dt} - C \frac{dT}{dt} \quad (2.5)$$

As the result of this complex function, a specific calibration procedure should be used. The calibration procedure has two identical constant heating rate measurements; first experiment should be performed without sample and pan (with empty furnace), and second experiment is carried out with sapphire disks which are placed on the sample and reference positions without pans and lids. Such experiments compute the resistance  $R_{SCP}$  and  $R_{RCP}$ , and capacitance  $C_{SCP}$  and  $C_{RCP}$  of the cell. Then, the temperature and energy calibrations are carried out using indium standard ( $T_m = 156.6$  °C and  $\Delta H_m = 28.6$  J.g<sup>-1</sup>). Benzophenone can also be used as a second standard for the cell calibration in temperature ( $T_{m, benzophenone} = 48.0$  °C). The calibrations of temperature and energy have to be repeated according to the experimental scanning rate. In order to provide a good signal-to-noise ratio, the sample masses of 5-10 mg were used. Scanning rate was selected as 10 K.min<sup>-1</sup> to ensure the good resolution of examined thermal events. All experiments were carried out under a nitrogen atmosphere.

### 2.2.3 Modulated temperature DSC (MT-DSC)

In standard DSC, sample undergoes the thermal events which affect its physical and/or chemical properties during the measurements. These thermal events consist of the glass transition, crystallization, and melting, as well as curing and evaporation. The “reversing” and “non-reversing” events may occur at the same time or in the same range of temperature. Thus, the average heat flows are overlapped and cannot be differentiated using standard DSC curves. In 1993, Reading et al. [13] proposed to express the heat flow measured from standard DSC as follows:

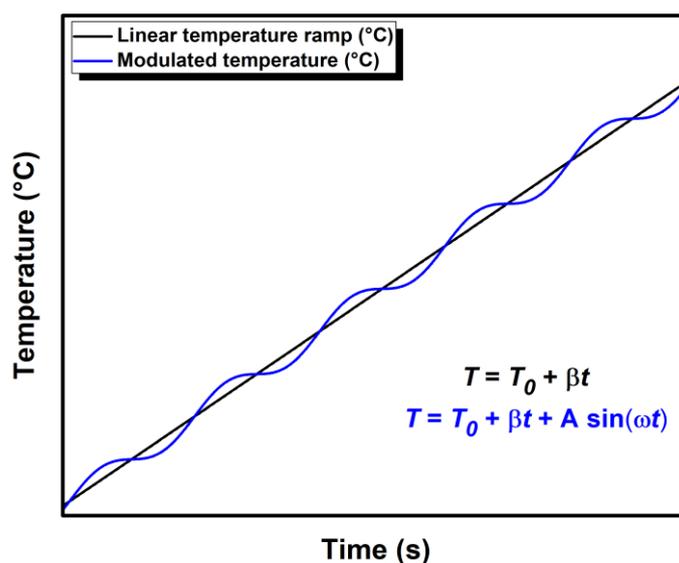
$$\Phi = \frac{dQ}{dt} = C^* \beta + f(t, T) \quad (2.6)$$

where  $f(t, T)$  corresponds to the part of the heat flow related to the kinetic or non-reversing events, so called non-reversing heat flow  $\Phi_{NR}$ . The reversing heat flow  $\Phi_R$  linked to the

thermodynamic events corresponds to the complex heat capacity and the scanning rate  $C^*\beta$ . The main difference of MT-DSC from standard DSC is the control of the sample temperature and the data treatment method. In addition to the linear temperature ramp as in standard DSC, MT-DSC includes a temperature modulation program in which the sample temperature follows a periodic wave pattern (such as a sinusoidal wave, shown in Fig. 2.7.). Reading et al. [14] proposed to super-impose sinusoidal oscillation with the linear temperature ramp in order to separate the reversing and non-reversing events:

$$T = T_i + \beta t + A \sin(\omega t) \quad (2.7)$$

where  $A$  is the modulation amplitude and  $\omega$  is the angular frequency of the temperature modulation with the period of oscillation  $P = 2\pi/\omega$ .



**Figure 2. 7.** Illustration of heating ramp in the MT-DSC experiment on PDLA sample with a modulation amplitude  $A = \pm 0.318$  K, a period of modulation  $P = 60$  s ( $\omega = 2\pi/P$ ) and underlying heating rate  $\beta_h = 2$  K.min<sup>-1</sup>. These experimental parameters correspond to a “heat-cool” mode.

The derivative of a sinusoidal temperature with respect to time gives the modulated heat flow  $\Phi$  as follows:

$$\Phi = \frac{dQ}{dT} = C^* (\beta + A\omega \cos(\omega t)) \quad (2.8)$$

and the complex heat capacity  $C^*$  is defined as:

$$C^* = \frac{A_{HF}}{A_\beta} = \frac{\omega AC^*}{\omega A} \quad (2.9)$$

where  $A_{HF}$  and  $A_\beta$  are the amplitude of the heat flow modulation and of the heating rate modulation, respectively.

The thermodynamic events linked to the vibrational and translational motions of molecules can be investigated from the reversing heat flow  $\Phi_R$ . Such motions are very rapid and can instantaneously follow any sample temperature modulation. On the contrary, the non-reversing heat flow  $\Phi_{NR}$  cannot follow the modulation and contribute to the modulated part of the heat flow [15]. Thus, the total DSC signal simultaneously measured by MT-DSC corresponds to the signal measured by standard DSC and the reversing DSC. In that case, the reversing heat flow  $\Phi_R$  and the non-reversing heat flow  $\Phi_{NR}$  can be summarized according to:

$$\Phi_R = C^* \beta = \frac{A_{HF}}{A_\beta} \beta \quad (2.10)$$

$$\Phi_{NR} = \frac{dQ}{dt} - C^* \beta \quad (2.11)$$

A phase lag  $\varphi$  consists of the total heat flow and the heating modulation. In this situation, two evident heat capacity components considered as the in-phase component related to  $\Phi_R$   $C'$  and the out-of-phase component related to  $\Phi_{NR}$   $C''$  can be determined by following equations:

$$C' = |C^*| \cos(\Phi) \quad (2.12)$$

$$C'' = |C^*| \sin(\Phi) \quad (2.13)$$

The MT-DSC measurements were carried out with the Q100 DSC from Thermal Analysis Instruments (DSC Q100 TA Instruments). All experiments were performed under a nitrogen atmosphere and the calibration of MT-DSC was carried out in three different steps:

1. the heat flow, the temperature and the baseline were calibrated by using  $T_{zero}^{TM}$  technology;
2. the temperature and the energy calibration were performed by using indium;

3. the specific heat capacity calibration was carried out using sapphire as a standard as it did not undergo any transition in the studied temperature range.

The heat capacity of sapphire as a function of temperature is known exactly and is very stable [16]. The third calibration step has to be repeated if the amplitude and period of oscillation, the average heating or cooling rate are changed. In the end, a calibration factor  $K_{CP}$  is calculated from the ratio of the experimental and theoretical specific heat capacity values of sapphire:

$$K_{CP}(T) = \frac{C_{P,Sapphire,theoretical}(T)}{C_{P,Sapphire,experimental}(T)} \quad (2.14)$$

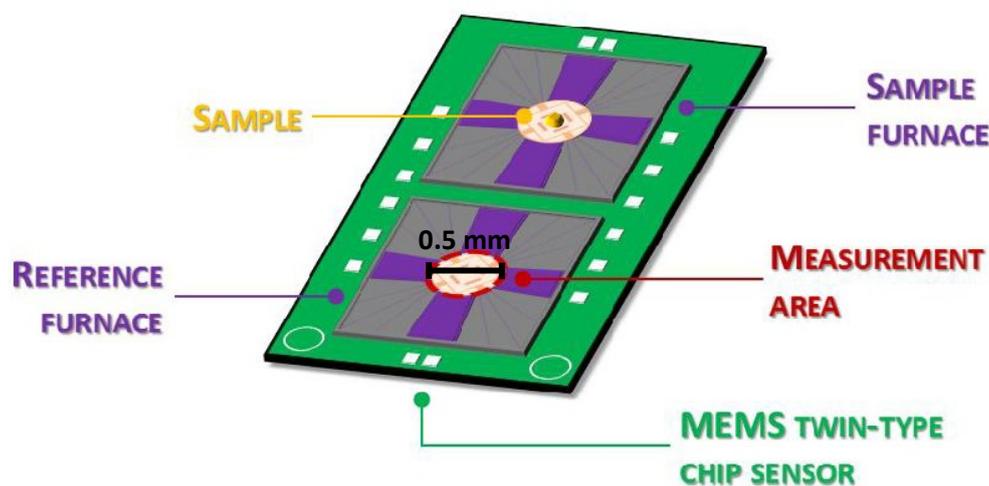
This factor is averaged on the temperature range of measurements and can be then used to correct the apparent heat capacity of sample.

In this work, MT-DSC analysis was performed in heat-cool mode to investigate the glass transition evolution of molecular dynamics in terms of free volume and CRR as well as to perform crystallinity degree calculation. The measurements were carried out with a modulation amplitude of  $\pm 0.318$  K, a period of 60 s and a scanning rate of 2 K.min<sup>-1</sup>.

#### **2.2.4 Fast scanning calorimetry (FSC)**

Some thermal events or reactions may take place simultaneously during heating. While MT-DSC analysis allows us to dissociate the reversing and non-reversing events, Schick and Mathot proposed to apply very high scanning rates in order to avoid or prevent thermal reactions, as well as to observe very fast thermal events and/or crystallization processes [17]. In this case fast scanning calorimetry (FSC) is used. The fundamental principle of FSC is based on the measurement of the difference in amount of required heat to increase the temperature of a sample and a reference. The investigated sample and the reference are placed in separated furnaces. Then, both furnaces are heated and heat is transferred to the sample and the reference. The difference in thermal power which required to maintain the sample and the reference at the same temperature is monitored. In this work, FSC analysis was performed by fast scanning calorimeter Flash DSC 1 from Mettler Toledo with the power compensation twin-type calorimetric chip sensor, based on micro-electro-mechanical systems (MEMS)

technology. As shown in Fig. 2. 8, the twin-type chip sensor has two independent furnaces – one for a sample and the other one for reference. The MEMS chip sensor is made by a stable ceramic and the applied temperature is transferred by electrical thermocouples.



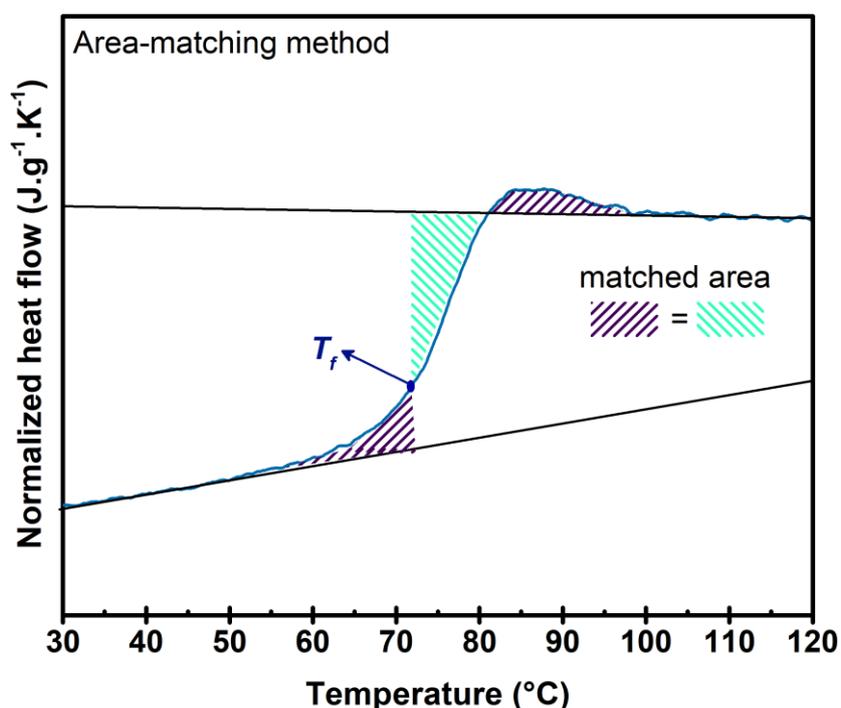
**Figure 2. 8.** Illustration of the twin-type chip sensor based on MEMS technology [18].

Both sides of the chip sensor contain two thermal resistance heaters in order to apply desired temperature and the temperature resolution is calculated by the time constant of the sensor (about 1 millisecond, i.e. thousand times less than in case of standard DSC). The heat flow is controlled through eight thermocouples at each side (totally sixteen thermocouples). The measurement area of the chip is fabricated of a silicon nitride and silicon dioxide coated with a thin layer of aluminum to provide homogenous temperature distribution. Before each use the chip sensor is conditioned and corrected according to Mettler Toledo procedures. The conditioning procedure allows verify the proper behavior of the sensor and its potential memory effects are erased by sensor heating to the maximum temperature (400 °C). The correction procedure is carried out in order to correct the thermocouple signal with respect to the sensor support temperature. The correction procedure is performed by comparing the signal of thermocouples with the temperature behavior of the heating resistance. In this work, small piece of sample was cut and placed directly in the center of the measurement area of a MultiSTAR UFS 1 MEMS chip sensor. A Huber TC100 intra cooler was used to cool the system down to -90 °C and carry out high cooling rates. The calorimeter was operated under nitrogen

gas flow of  $20 \text{ mL}\cdot\text{min}^{-1}$  to avoid water condensation from the environment and to optimize the applied program temperature [19]. The mass of samples was estimated from the amplitude of the heat capacity step at the glass transition by comparing the value of the heat capacity step ( $\Delta C_p$ ) obtained by standard DSC at  $|\beta_c| = \beta_h = 0.78 \text{ K}\cdot\text{s}^{-1}$  and the value of  $\Delta C_p$  obtained by FSC at  $|\beta_c| = \beta_h = 1500 \text{ K}\cdot\text{s}^{-1}$  [18]. The heat flow curves hereby were normalized to the mass and the scanning rate, and the spectra curves were then plotted in equivalent  $C_p$  unit, i.e.  $\text{J}\cdot\text{g}^{-1}\cdot\text{K}^{-1}$ .

#### 2.2.4.1 Glass transition temperature determination by the fictive temperature concept

The respective glass transitions have been investigated in terms of fictive temperature and as a function of cooling rate. Thermal history of homopolymers and sc PLA samples was erased at  $220 \text{ }^\circ\text{C}$  and  $270 \text{ }^\circ\text{C}$ , respectively, by isotherms during  $0.5 \text{ s}$  before any investigations. Samples were then studied at temperatures from  $-50 \text{ }^\circ\text{C}$  to  $270 \text{ }^\circ\text{C}$  at scanning rate  $|\beta_c| = \beta_h$  ranging from  $300 \text{ K}\cdot\text{s}^{-1}$  to  $4000 \text{ K}\cdot\text{s}^{-1}$  to promote the resolution of investigated thermal events. Fig 2.9 presents the normalized heat flow of amorphous sc PLA sample as a function of temperature in the glass transition region.



**Figure 2. 9.** FSC normalized heat flow curve of wholly amorphous sc PLA sample as a function of temperature upon heating at scanning rate  $|\beta_c| = \beta_h = 1500 \text{ K}\cdot\text{s}^{-1}$ .

The values of the glass transition temperature were then calculated as the fictive temperatures by using area matching method that was proposed by Moynihan et al. [20], as shown in Fig 2.9. Since the heat capacity step at the glass transition is independent on the scanning rate, special attention is paid to the sensitivity, i.e. signal-to-noise ratio, of the heat flows during FSC experiments because of its dependence on the sample mass and scanning rate [21]. The cooling rate dependence of the fictive temperature is extensively presented in literature [22]. When applied cooling rate increases both the glass transition temperature and fictive temperature will increase according to the non-Arrhenius behavior. Such cooling rate dependence of the fictive temperature can be fitted by the Vogel-Fulcher-Tamman-Hesse (VFTH) equation [23]:

$$\log(|\beta_c|) = A_\beta - \frac{B}{T_f - T_V} \quad (2.15)$$

where  $A_\beta$  is a constant,  $B$  is a fitting parameter and  $T_V$  is a Vogel temperature which is the extrapolated temperature when relaxation time  $\tau$  approaches infinite. In the concept of the fictive temperature, the VFTH equation can be modified to the Vogel-Fulcher-Tamman (VFT) equation to determine the dynamic fragility index  $m$  as follows [24, 25]:

$$m = \left[ \frac{d(\log|\beta_c|)}{d\left(\frac{T_g}{T}\right)} \right]_{T=T_g} \quad (2.16)$$

Taking into account the VFTH behavior for the cooling rate dependence of the fictive temperature,  $m$  can also be expressed as follows:

$$m = \frac{B.T_g}{(T_g - T_V)^2} \quad (2.17)$$

The fictive temperatures as a function of cooling rate were determined by area-matching method for all the samples and the experimental data were fitted by the VFTH function to calculate the fragility index  $m$ .

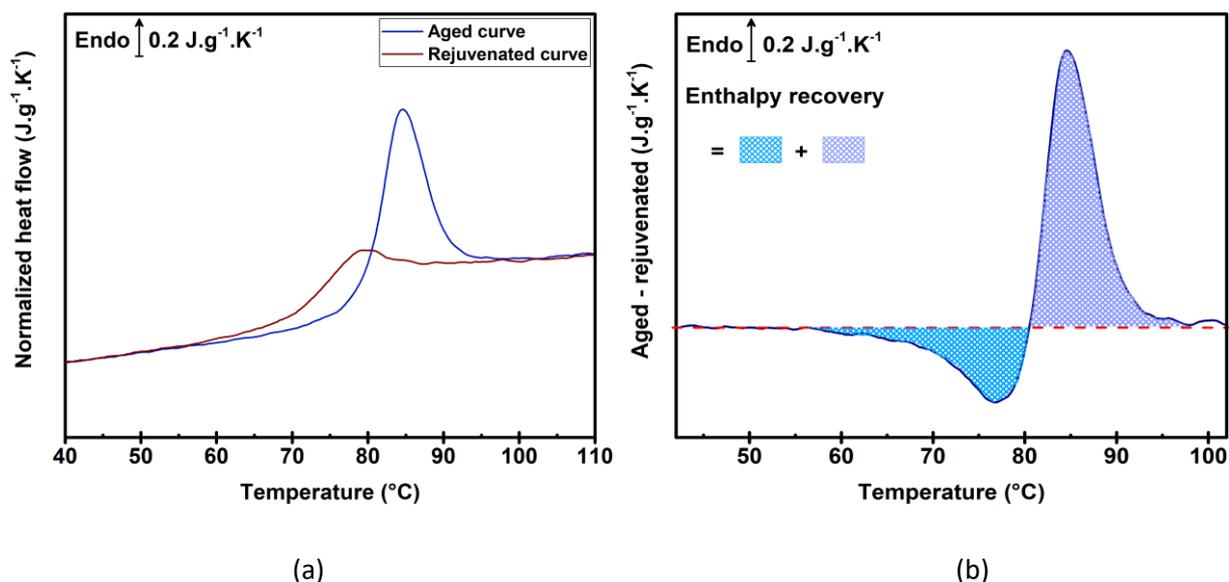
#### **2.2.4.2 Thermal lag corrections**

In order to improve the accuracy of the fictive temperatures investigation by FSC, the thermal lag correction is required [17]. Thermal lag ( $\delta T_L$ ) is defined as a shift in temperature due to the sample thickness and the high scanning rate, which provokes the heat transfer delay between heater and sample [26]. The thermal lags can be corrected by the melting of indium piece placed on the top of the sample [26-29]. As proposed by Schawe [27] two corrections of thermal lag may be applied: the static ( $\delta T_S$ ) and the dynamic ( $\delta T_D$ ) thermal lag. The static thermal lag ( $\delta T_S$ ) corresponds to a third of the difference between the onsets melting point of an indium piece placed on the sample specimen and another one placed directly on the reference chip sensor, when extrapolated to heating rate zero. The dynamic thermal lag ( $\delta T_D$ ) corresponds to a half of the difference between the fictive temperatures determined from measurements upon heating and cooling. A total thermal lag values were then calculated for all samples by following equation:

$$\delta T_L = \delta T_S + \delta T_D \quad (2.18)$$

#### **2.2.4.3 Physical aging in a scanning rate range more than six decades**

Physical aging was carried out at different aging temperatures ( $T_{ag} = T_g - 18 \text{ }^\circ\text{C}$ ) calculated according to the corrected  $T_g$  for different aging time  $t_{ag} = 1 \text{ min}; 10 \text{ min}; 100 \text{ min}$  at the scanning rate  $|\beta_c| = \beta_h$  ranging from  $300 \text{ K}\cdot\text{s}^{-1}$  to  $4000 \text{ K}\cdot\text{s}^{-1}$ . Fig 2. 10a displays the normalized heat flows of aged and rejuvenated sample. Fig 2. 10b shows the scheme of enthalpy recovery determination of a glass, aged at a temperature  $T_{ag}$  during an aging time  $t_{ag}$  by the equations 1.23 and 1.24 [30].



**Figure 2. 10.** (a) Heat flows of aged and rejuvenated (non-aged) curves of amorphous sc PLA sample. (b) Schematic illustration of the enthalpy recovery calculation.

### 2.2.5 Dielectric relaxation spectroscopy (DRS)

DRS is a powerful technique to investigate molecular dynamic and relaxation phenomena of materials, especially polymers. The advantage of DRS is that measurements can be carried out at 9 decades of frequency or time, and this range can be extended up to 12 decades [31]. DRS analysis is based on the study of the motion of permanent dipoles present in a material as an impact of applied alternating electric field  $E(\omega)$ . In case of polymer, the net dipole moment per unit volume is linked to the total vector of all molecule dipoles that exist in the repeating unit, the polymer chain and overall structure in polymer chain. The electric field stimulates the electronic cell according to the atomic nucleus (so called electronic polarization) and composes induced dipole moment in atoms (so called atomic polarization). Such phenomenon in polymers is due to the rotational mobility of permanent dipole moments ( $\mu$ ), which are defined as dipole moments of chemical groups in the polymer chains. This reorientation of the permanent dipoles of the molecules allows us to determine the sample properties, i.e. permittivity, energy storage, resistivity and dissipation, and, thus, to investigate molecular motions occur within the polymers as a function of frequency and temperature.

Therefore, dielectric relaxation spectroscopy technique is usually used to investigate the molecular mobility [32-35]. In the frequency range of  $10^{-6}$  to  $10^7$  Hz, the sample can be studied as a circuit that is composed of an ideal capacitor and an ohmic resistor combined in parallel or serial. The complex impedance  $Z^*(\omega)$  of the circuit is measured by the spectrometer and may be defined in terms of energy dissipation or resistance  $R(\omega)$ , and energy storage or capacitance  $C(\omega)$ , in which  $\omega$  the angular frequency is determined from the frequency  $f$  by the following equation:

$$\omega = 2\pi f \quad (2.19)$$

The derivation of the complex electrical impedance  $Z^*(\omega)$  gives other properties, such as electrical modulus  $M^*(\omega)$ , electrical conductivity  $\sigma^*(\omega)$ , resistivity  $R^*(\omega)$ , and especially complex dielectric permittivity  $\varepsilon^*(\omega)$ . To measure complex impedance  $Z^*(\omega)$ , a sinusoidal voltage  $U^*(\omega)$  at a constant frequency is applied to the sample:

$$U^*(\omega) = U_0 \exp(j(\omega t)) \quad (2.20)$$

then, the current  $I_s^*(\omega)$  is measured as follows:

$$I_s^*(\omega) = I_0 \exp(j(\omega t + \varphi)) \quad (2.21)$$

where  $\varphi$  is the phase shift between the applied voltage and measured current. The complex impedance  $Z^*(\omega)$  is then calculated by following ratio [36]:

$$Z^*(\omega) = \frac{U^*(\omega)}{I^*(\omega)} \quad (2.22)$$

As shown in Fig.2.11, the impedance is determined from the measurements of two voltages corresponding to the generated voltage applied to the sample and the voltage-converted sample current  $I_s(\omega)$ . These voltages are analyzed by the Fourier transform technique to obtain information on their phases and amplitudes. The complex dielectric permittivity ( $\varepsilon^*$ ) of a capacitor  $C^*$  filled with studied sample is expressed as:

$$\varepsilon^*(\omega) = \frac{C^*(\omega)}{C_0} \quad (2.23)$$

where  $C_0$  is the capacitance of the empty capacitor. The capacitance of the empty capacitor can be defined as [36]:

$$C_0 = \frac{\varepsilon_0}{d} A \quad (2.24)$$

where  $\varepsilon_0$  is the permittivity of vacuum,  $A$  is the area of the electrode and  $d$  is the thickness of the sample (distance between the plates as shown in Fig. 2.11). The complex dielectric permittivity ( $\varepsilon^*$ ) can be derived by measuring the complex impedance  $Z^*(\omega)$  of the sample:

$$\varepsilon^*(\omega) = \frac{1}{i\omega Z^*(\omega)C_0} \quad (2.25)$$

$$\varepsilon^*(\omega) = \varepsilon'(\omega) - i\varepsilon''(\omega) \quad (2.26)$$

where  $\varepsilon'(\omega)$  is the real part (related to the stored energy) and  $\varepsilon''(\omega)$  is the imaginary part (related to the dissipation of energy) of the complex dielectric permittivity ( $\varepsilon^*$ ).

Permittivity allows us to investigate the ability of charges in a material to be reoriented or displaced in the presence of an external electric field in order to obtain information about the capacity of dipoles in a material mobility. In this study, DRS measurements were carried out with a Novocontrol Alpha analyzer. The temperature was controlled by a Quatro Novocontrol Cryosystem with temperature stability better than  $\pm 0.2$  K. Nonmetalized samples were placed between parallel 30 mm diameter inox plated electrodes. The broadband dielectric converter (Alpha analyzer interface) allows the measurement of the complex dielectric permittivity (real and imaginary parts) in a frequency range from  $10^{-1}$  Hz to  $10^6$  Hz. The temperature was changed between  $-150$  °C and  $150$  °C with increasing step of  $10$  °C for the range from  $-150$  °C to  $50$  °C, and  $1$  °C for the range from  $50$  °C to  $150$  °C. During the whole measurements, the samples were kept in a pure nitrogen atmosphere.

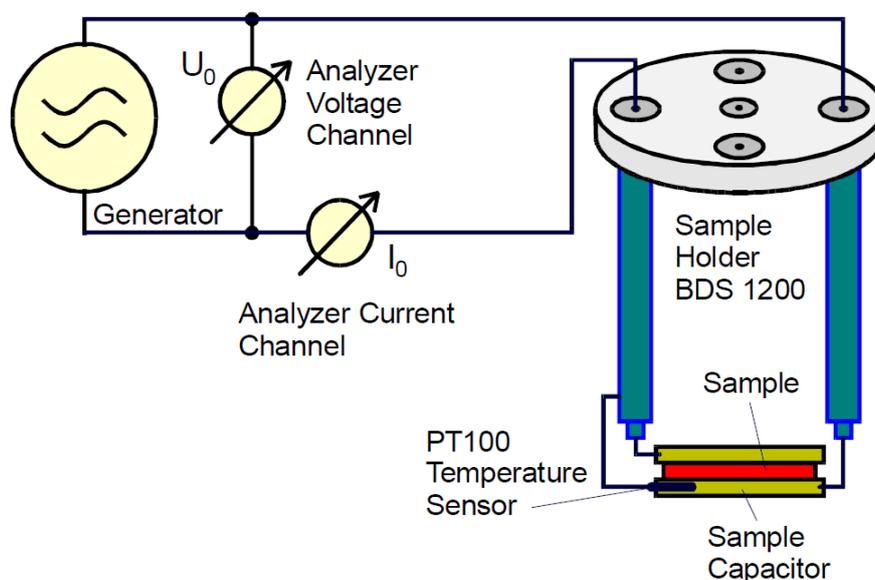


Figure 2. 11. Schematic representation of the spectrometer analyzer [37].

### 2.2.6 X-ray diffraction (XRD)

XRD is an analytical technique used for phase identification of crystalline phases. This analysis is based on a constructive interference of monochromatic X-rays and a crystalline sample and provides information on unit cell dimensions. X-rays are generated in a cathode ray tube by heating a filament to produce electrons, accelerating electrons towards a target material by applying a voltage, and bombarding the target material with electrons. When electrons have sufficient energy to dislodge inner shell electrons of the target material, characteristic X-ray spectra are obtained. As shown in Fig. 2.12, the basic principle of X-ray diffraction is based on the bombarding of a sample with an X-ray beam and the recording the intensity of the reflected X-rays. Constructive interference and peak intensity occur when the geometry of the loaded X-rays impinging the sample follows the Bragg's law [38]:

$$n\lambda = 2d \sin(\theta) \quad (2.27)$$

where  $d$  is the spacing between diffracting planes in the crystalline lattice,  $\theta$  is the incident angle,  $n$  is any integer, and  $\lambda$  is the wavelength of the beam (depicted in Fig 2.13).

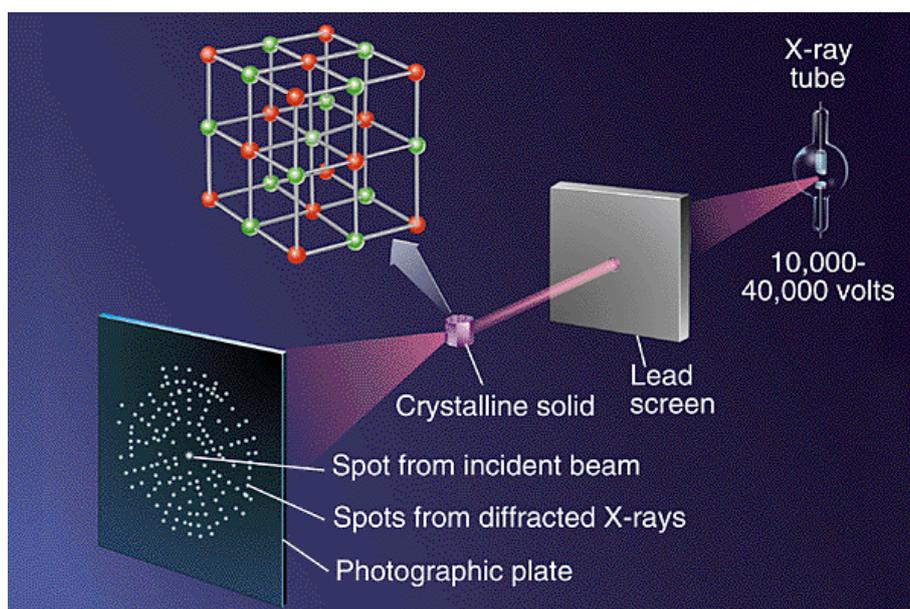


Figure 2. 12. Schematic representation of the basic principle of X-ray diffraction [39].

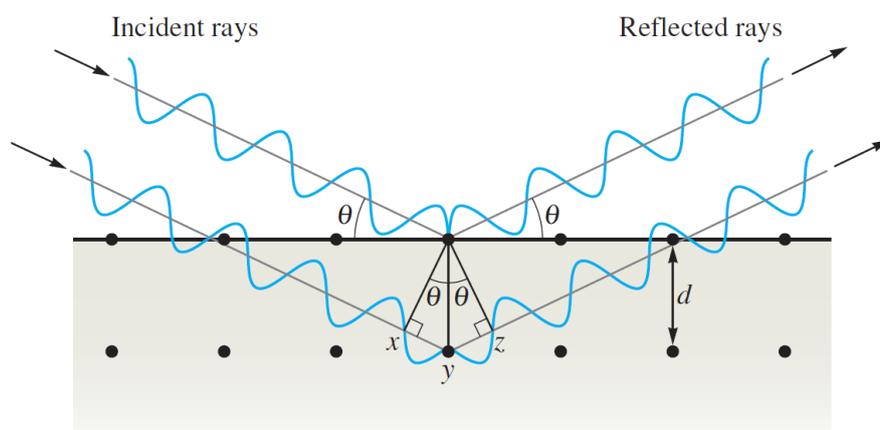


Figure 2. 13. Illustration of the Bragg's law [39].

X-ray diffraction technique is widely used for:

- ◆ characterization of crystalline materials;
- ◆ determination of unit cell dimensions;
- ◆ measurement of sample impurity;
- ◆ characterization of thin films by:
  - determining lattice mismatch between film and substrate;
  - determining dislocation density and quality of film;
  - determining the thickness, roughness and density.

In this work, XRD measurements were performed on Bruker's X-ray Diffraction D8-Discover instrument. The generator was set up at 35 kV and 40 mA and the target material copper for single-crystal diffraction with  $\text{CuK}\alpha$  radiation ( $\lambda = 1.54 \text{ \AA}$ ) was selected. The measurements were run at room temperature ( $T = 25 \pm 2 \text{ }^\circ\text{C}$ ) from  $5^\circ$  to  $40^\circ$  at  $0.04^\circ$  increment with 1s/step scan speed. All X-ray patterns were corrected by background scattering.

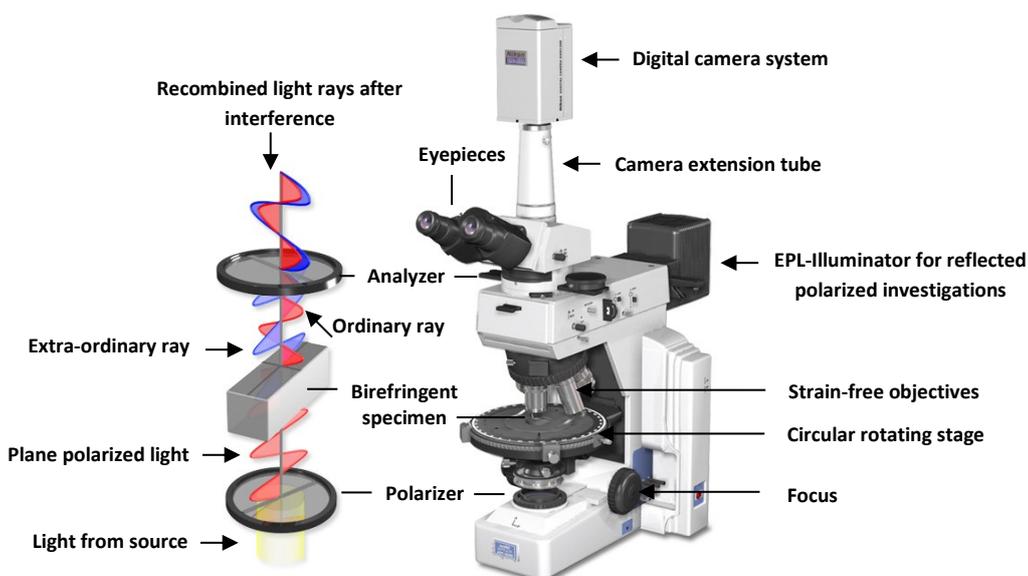
### **2.2.7 Polarized optical microscopy (POM)**

POM is a basic and sensitive technique used for both quantitative and qualitative studies of a wide range of anisotropic specimens. POM analysis is designed to observe and photograph specimens that are visible primarily due to their optically anisotropic character. In order to accomplish this task, POM uses polarized light. Therefore, in experiment two additional polarizing filters are associated to a conventional optical microscope:

- polarizer, positioned in the light path before the specimen;
- analyzer, placed in the optical pathway between the objective rear aperture and the observation tubes or camera port.

Such filters are used to restrict electro-magnetic field vectors constituting visible light to a single plane, as shown in Fig. 2. 14. As a consequence, all electro-magnetic field vectors are vibrating in the same plane and the light is then polarized.

In cross polarization mode, the analyzer is oriented perpendicularly to the first polarizer. This position leads to block remaining electro-magnetic field vectors. As a result, while no sample is placed on the light field, the light polarized by the polarizer is blocked by the analyzer, resulting in no visible light in the eyepiece. However, when birefringent sample, such as anisotropic material, is placed on the polarized light field, the restricted electro-magnetic field vectors coming through the sample are produced in many different perpendicular planes due to the refractive index depending on the polarization. Then, electro-magnetic field vectors which are parallel to the polarization direction of the analyzer can pass through and are visualized in the eyepiece.

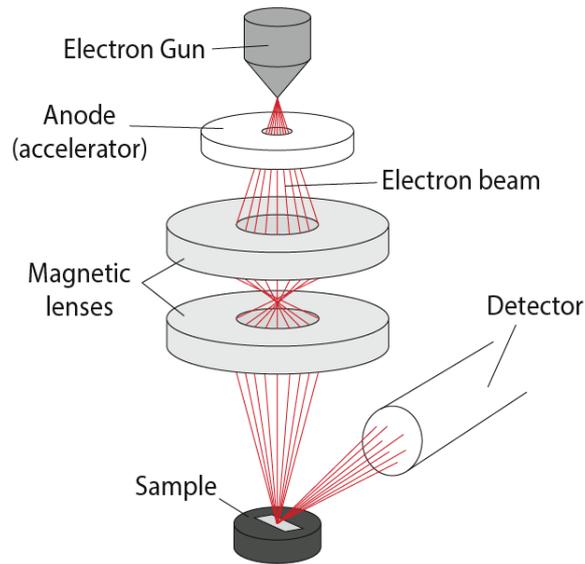


**Figure 2. 14.** Schematic representation of polarized optical microscopy configuration [40].

In this work, POM observations were carried out by using a universal Nikon EPI-illuminator with Nikon M Plan x2.5 / x5 / x10 lenses connected with a digital sight camera system from Nikon Corporation, either transmission or reflection mode. A Mettler FP82HT hot stage connected to a Mettler FP90 central processor was combined with POM as a temperature controller system during the crystallization observations.

### 2.2.8 Scanning electron microscopy (SEM)

During SEM analysis a sample is placed in a vacuum chamber and its surface is scanned by a focused beam of electrons. Due to the particle-wave duality of electrons, the electron beam is accelerated and focused on a sample in the path of anode, condenser and magnetic lenses, as shown in Fig 2. 15. These electrons interact with atoms in the sample as well as the sample surface. The primary beam focused on the sample area leads to the ejection of other electrons including secondary electrons, back-scattered electrons or X-rays. Specific detectors are used according to the materials nature and the signals measured involve information about the surface topography and the composition of the sample. In this work, SEM analysis was carried out by a Dual-BEAM LEO1530-ZEISS scanning electron microscopy to observe the surface morphology of semi-crystalline homopolymers PLLA and PDLA, and sc PLA.



**Figure 2. 15.** Schematic representation of scanning electron microscopy analysis in a vacuum chamber.

## 2.2.9 Permeation measurements

### 2.2.9.1 Water permeation measurements

The permeation process includes several steps (shown in Fig 2.16) such as sorption, diffusion and desorption while maintaining the concentration of the permeant as a constant at the upstream interface of the film ( $C'$ ). The concentration of the permeant at the downstream interface of the film ( $C''$ ) is negligible compared to  $C'$  ( $C' \gg C''$ ). In case of this concentration difference, a transfer of matter is observed and follows the two laws of Fick [41]. The dissolution-diffusion process results from the association of two phenomena:

- the dissolution of permeant molecules within the material which is characterized thermodynamically by a solubility coefficient ( $S$ );
- the diffusion of permeant molecules through the material (kinetic order) and characterizes by diffusion coefficient ( $D$ ).

The first law of Fick explains the dependence of the flux  $J(x, t)$  on the perpendicular position to the plane of the film ( $x$ ) and the time ( $t$ ) with the gradient of the local concentration  $C(x, t)$ :

$$J(x, t) = -D \frac{\partial J(x, t)}{\partial x} \quad (2.28)$$

where  $D$  is the local diffusion coefficient of permeant. According to the observed Fickian mechanism [42] two types of diffusion can be present. The type I is observed when  $D$  is constant and the type II occurs when  $D$  depends on the concentration of the permeant. According to Fick's second law, the evolution of the local concentration with the divergence of the flux can be expressed as follows:

$$\frac{\partial C(x,t)}{\partial t} = \frac{\partial J(x,t)}{\partial x} \quad (2.29)$$

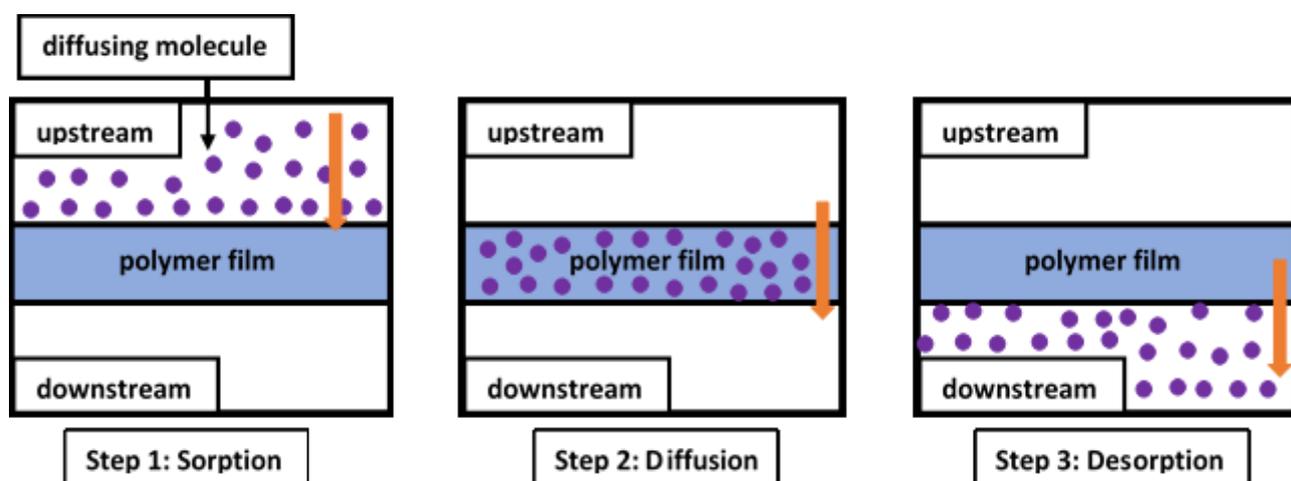


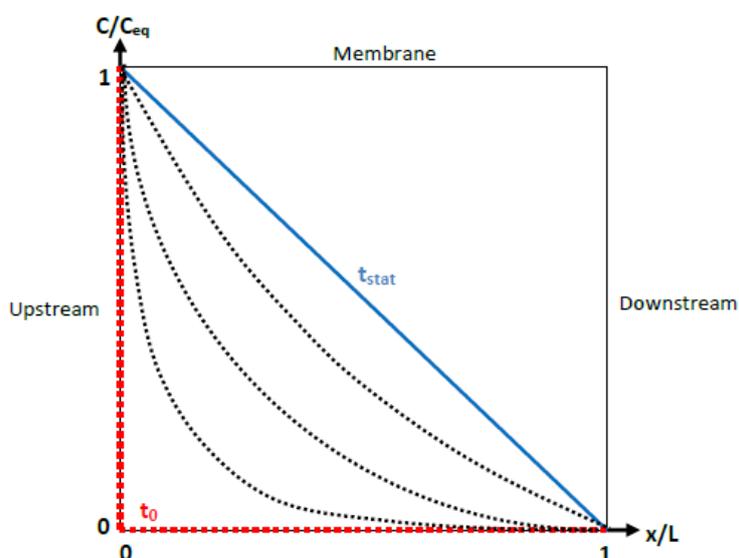
Figure 2. 16. Schematic representation of the principle of permeation measurements in 3 steps.

The interfaces of the film can be defined as  $x = 0$  for upstream and  $x = L$  for downstream, where  $L$  corresponds to the film thickness. Several hypotheses must be made in order to apply these diffusion models: (1) the sample is homogenous and initially no trace of permeant is present, (2) the impact of swelling during measurements is negligible, and (3) the sorption at the film surface reaches equilibrium in a quasi-instantaneous manner. The initial and limiting conditions are expressed by following functions and the permeant concentration profiles are depicted schematically in Fig 2.17.

$$\text{when } t = 0 \quad 0 < x < L \quad C(x, 0) \quad (2.30)$$

$$\text{when } t > 0 \quad x = 0 \quad C(0, t) = C' = C_{eq} \quad (2.31)$$

$$\text{when } t > 0 \quad x = L \quad C(L, t) = C'' \approx 0 \quad (2.32)$$



**Figure 2. 17.** Schematic representation of the profiles of permeant concentration in reduced scales at  $t = 0$  up to steady state where  $t = t_{\text{stat}}$  for constant  $D$ .

The transport properties of the samples (semi-crystalline PLLA, PDLA, and sc PLA) towards water at 25 °C, 45 °C and 65 °C were studied to investigate the influence of stereocomplexation on film barrier properties. The water permeation measurements were carried out by the device designed by MPBM team of PBS laboratory (Fig 2.18). The permeation cell, in which the sample is installed, is placed in a thermoregulated chamber (oven at  $T = 25$  °C, 45 °C and 65 °C). Before measurement, a purge step is carried out with a flow of dry gas (Technical Nitrogen upstream and BIP downstream, Air Products®) applied for a certain time (generally 18h). The downstream gas arrives to a mirror hygrometer (General Eastern, USA) and the dew point temperature  $T_R$  (Dew point) is measured. When this temperature is sufficiently low (around -70 °C at atmospheric pressure), the flow of dry gas in upstream is stopped and liquid distilled water is added in the upstream compartment at a time  $t = 0$ . Due to the difference in the concentration of the permeant between the two compartments, the water molecules' transfer will result in a rise in the amount of water in the downstream compartment and so in a  $T_R$  increase over time, as shown in Fig. 2. 19. The measurements were repeated between 3 and 5 times for each sample and each temperature for better accuracy.

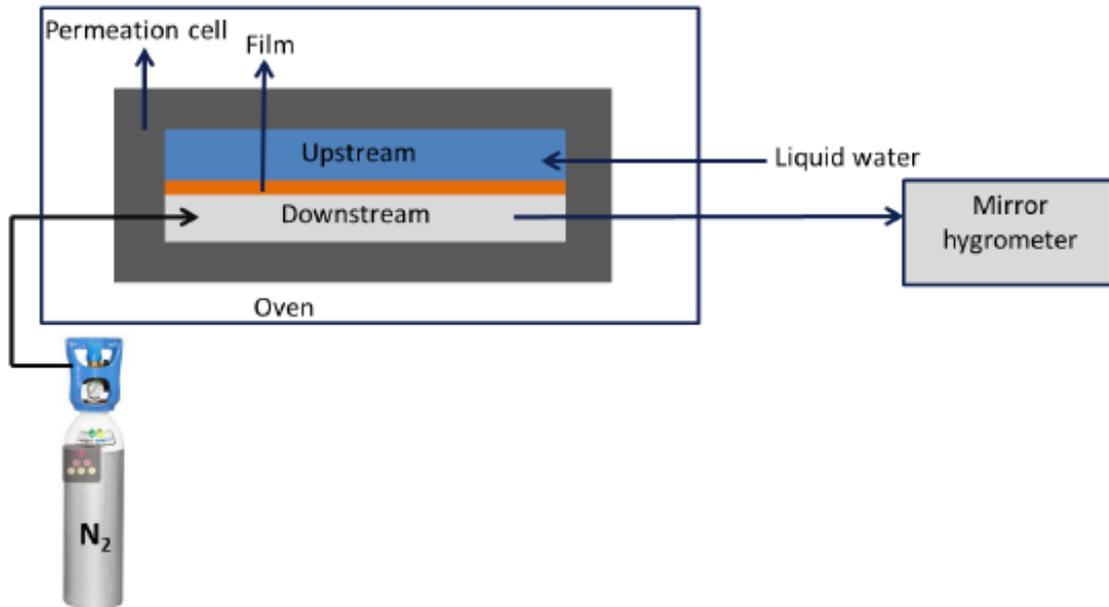


Figure 2. 18. Schematic illustration of the water permeation measurements.

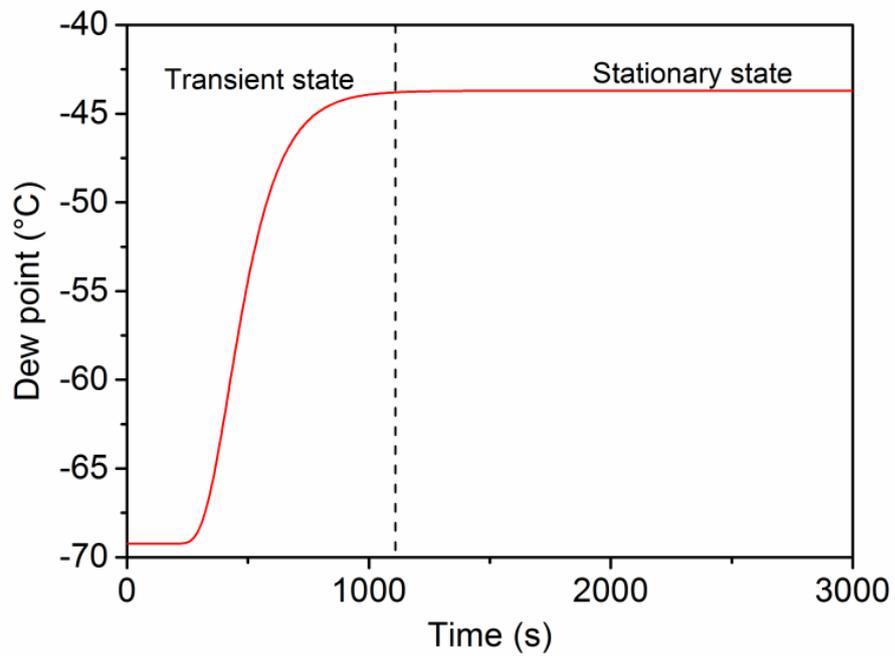


Figure 2. 19. Example of the evolution of dew point  $T_R$  over time during the water permeation measurement.

2.2.9.2 Gas permeation measurements ( $N_2$ ,  $O_2$ , and  $CO_2$ )

Gas permeability was measured by the so called “time-lag” method [43] by using the experimental device reported by Joly et al. [44] (shown in Fig. 2. 20). The gases used are: nitrogen  $N_2$  (Air Products®, purity 99.99%), oxygen  $O_2$  (Air Liquide®, purity 99.99%) and carbon dioxide  $CO_2$  (Air Products®, purity 99.5%).

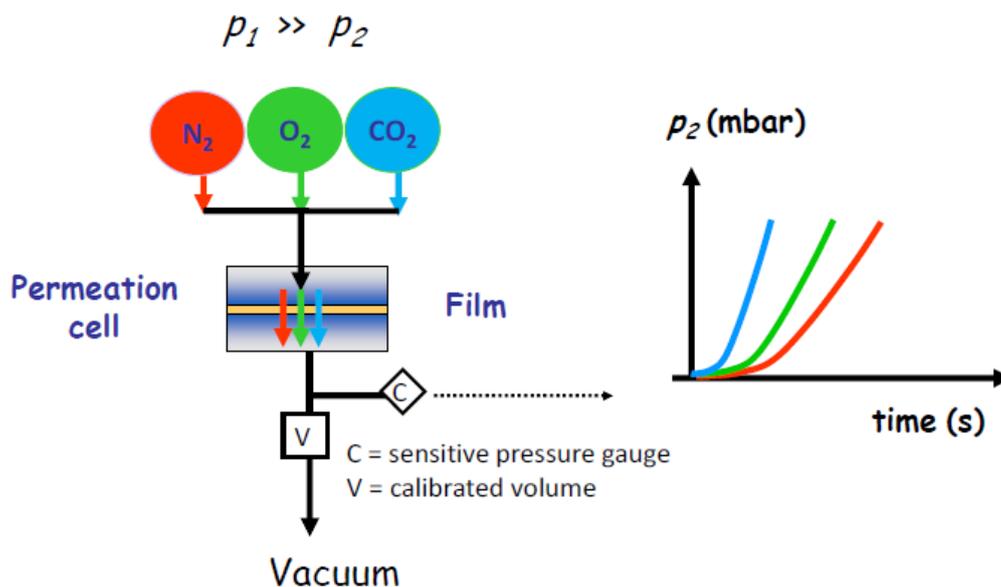


Figure 2. 20. Illustration of gas permeation apparatus.

Nitrogen, oxygen and carbon dioxide permeation properties of semi-crystalline PLLA, PDLA and sc PLA films at room temperature  $T = 25 \pm 2$  °C were determined using the permeation apparatus shown in Fig. 2. 20. Before measurements, the permeation cell (XX45047 Millipore filtration cell adapted for gas permeation) was completely evacuated by applying a vacuum on both sides of the film. Then, the upstream side was provided with the gas under test at pressure  $p_1$  (3 bar). The increase of pressure in the calibrated downstream volume  $p_2$  ( $p_1 \gg p_2$ ) was measured using a sensitive pressure gauge (0 – 10 mbar, Effa AW – 10 – T4) linked to a data acquisition system. The measurements of each gas were repeated between 3 and 5 times for each sample for better accuracy.



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***CHAPTER 3. INFLUENCE OF ELABORATION METHODS ON  
STEREOCOMPLEXATION REACTION***

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**CHAPTER 3. INFLUENCE OF ELABORATION METHODS ON STEREOCOMPLEXATION REACTION**

In order to study the stereocomplexation reaction, two different methods were used, i.e. the solvent casting and the extrusion. The experimental conditions (polymer concentration and homopolymer ratio in case of solution casting; and temperature and mixing time in case of extrusion) were optimized in terms of film microstructure and thermal properties.

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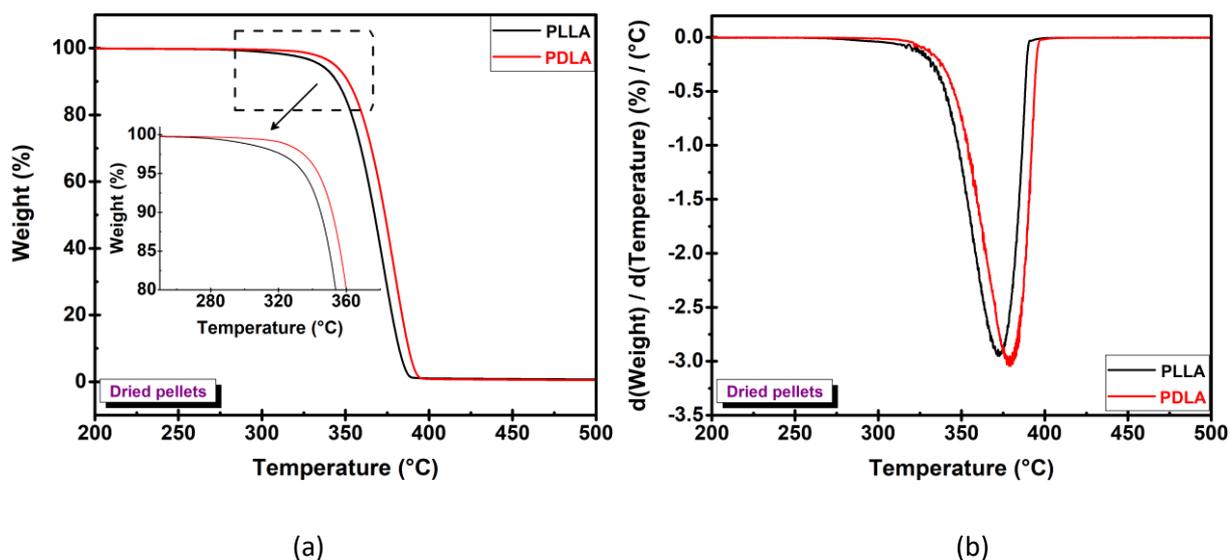
### 3.1 Homopolymers

#### 3.1.1 Homopolymer pellets

Dried PLLA and PDLA pellets were analyzed in order to investigate their thermal properties before further characterization.

##### 3.1.1.1 Thermal stability of pellets

Fig 3.1 presents thermograms and derivative curves of thermograms for PLLA and PDLA pellets as a function of temperature. Thermal degradation temperature ( $T_{deg}$ ) was determined as the temperature at which the mass loss was observed as shown in Table 3.1. PLLA pellets degrade about 25 °C lower than PDLA pellets ( $T_{deg} = 296$  °C and  $T_{deg} = 321$  °C, respectively) (Fig 3.1.a), while the maximum of derivative curves shows much less temperature variation between the homopolymers, i.e. about 5 °C (shown in Fig 3.1.b). In any case,  $T_{deg}$  of homopolymer pellets is sufficiently high for the stereocomplex formation.



**Figure 3. 1.** TGA curves of pure PLLA and PDLA pellets: (a) thermograms and (b) derivative curves as a function of temperature.

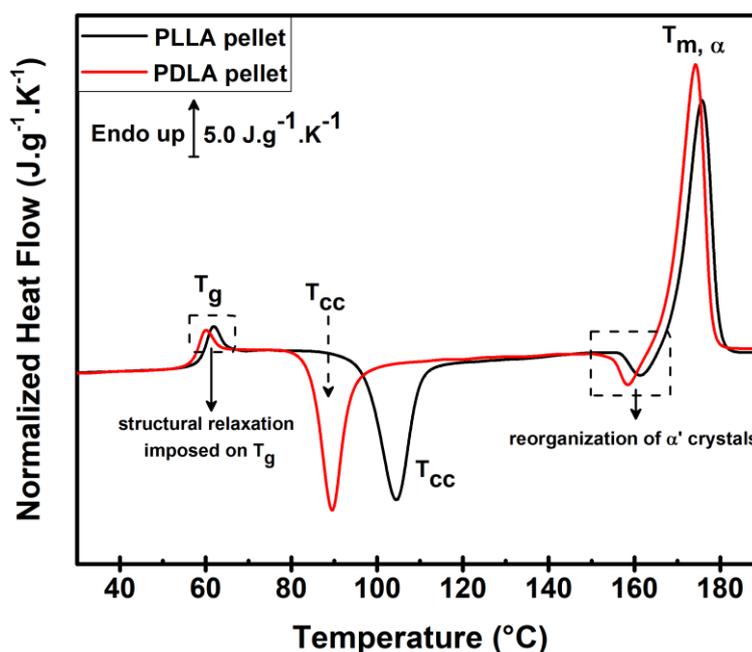
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Sample	$T$ of 1% weight loss (°C)	$T$ of 50% weight loss (°C)	Maximum of derivative curve (°C)
PLLA pellet	296	367	373
PDLA pellet	321	373	378

**Table 3. 1.** Degradation temperature ( $T_{deg}$ ) of pure PLLA and PDLA pellets at 1% and 50% weight loss and the maximum of derivative curves.

### 3.1.1.2 Thermal properties of pellets

DSC analysis was performed in order to examine the thermal properties of homopolymer pellets, i.e. glass transition temperature  $T_g$ , and melting temperature  $T_m$ . The DSC curves of homopolymer pellets are shown in Fig 3.2. The endothermic peaks at  $T_g$  ( $\sim 60$  °C) are due to the structural relaxation imposed by  $T_g$ . The endothermic peaks at  $\sim 175$  °C correspond to the melting of homo-crystals  $T_{m, \alpha}$  obtained by the cold-crystallization  $T_{cc}$  which is confirmed by the exothermic peaks at 104 °C and 90 °C for PLLA and PDLA, respectively.



**Figure 3. 2.** DSC curves of PLLA and PDLA pellets.

The cold-crystallization at temperature  $T < 120$  °C leads to  $\alpha'$  crystals form formation, i.e. disordered crystals with hexagonal packing [1, 2]. The reorganization of  $\alpha'$  imperfect crystals into  $\alpha$  form crystals is noticed by the exothermic peak before melting at temperature

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$T \approx 160$  °C. The values of the thermal parameters are presented in Table 3.2 and the crystallinity degree of  $\alpha$  crystals is estimated by following equation [3]:

$$X_c = \frac{[\Delta H_m - \Sigma \Delta H_{cc}]}{\Delta H_m^0} \quad (3.1)$$

where  $\Delta H_m$  is the measured melting enthalpy,  $\Sigma \Delta H_{cc}$  is sum of cold crystallization enthalpy,  $\Delta H_m^0$  is the melting enthalpy of 100 % crystalline sample, which equals 93 J/g for  $\alpha$  crystals of PLA [4]. The equal value of melting and cold-crystallization enthalpies reveals that homopolymer pellets are amorphous.

Sample	$T_g$ (°C)	$T_{cc}$ (°C)	$\Delta H_{cc}$ (J/g)	$T_m$ (°C)	$\Delta H_m$ (J/g)	$X_c$ (%)
<i>PLLA pellet</i>	60	104	47	176	48	1
<i>PDLA pellet</i>	59	90	51	174	53	2

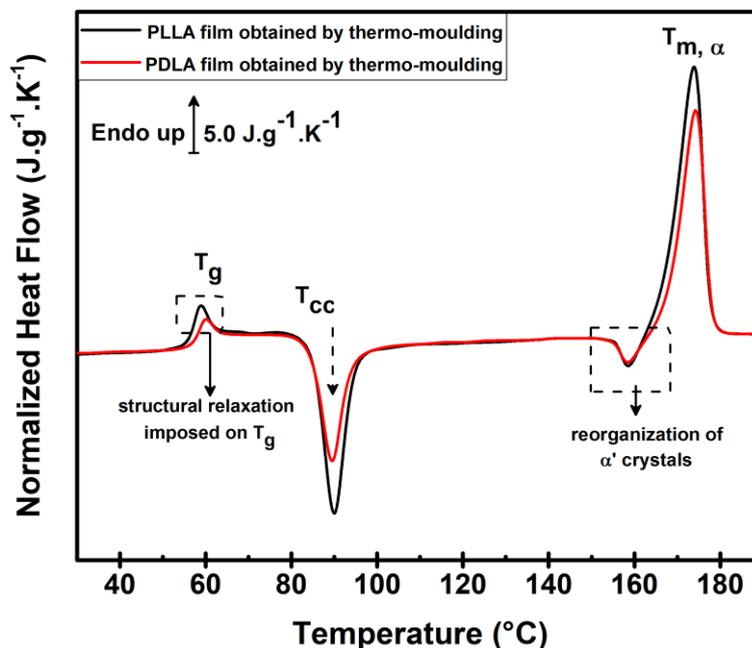
**Table 3. 2.** The values of the glass transition temperature  $T_g$ , the cold-crystallization  $T_{cc}$ , and the melting  $T_m$  and the enthalpy of the cold-crystallization ( $\Delta H_{cc}$ ) and the melting ( $\Delta H_m$ ) for PLLA and PDLA pellets.

### 3.1.2 Homopolymer film samples

Homopolymer films were obtained by thermo-molded and solution casting methods (presented in chapter 2). The influence of the elaboration methods on homopolymers properties was investigated by DSC, MT-DSC and XRD analysis.

#### 3.1.2.1 Amorphous homopolymers' film

Amorphous PLLA and PDLA films obtained by thermo-molding method were analyzed by DSC in order to examine thermal characteristics of homopolymers' films. Obtained DSC curves are presented in Fig 3.3. The values of the thermal parameters ( $T_g$ ,  $T_{cc}$  and  $T_m$ ) are presented in Table 3.3 and the crystallinity degree of  $\alpha$  crystals is estimated by Eq 3.1. DSC curves of the thermo-molded homopolymer films reveal the formation of  $\alpha'$  crystals that confirmed by the exothermic peaks at 90 °C and around 158 °C. The  $T_m$  of homopolymer crystals is observed at 174 °C for both PLLA and PDLA films. The close values of melting and cold-crystallization enthalpies reveal that the thermo-molded homopolymer films are amorphous.



**Figure 3. 3.** DSC curves of thermo-molded PLLA and PDLA films.

Sample	$T_g$ (°C)	$T_{cc}$ (°C)	$\Delta H_{cc}$ (J/g)	$T_m$ (°C)	$\Delta H_m$ (J/g)	$X_c$ (%)
<i>PLLA film</i>	58	90	51	174	53	2
<i>PDLA film</i>	59	90	43	174	45	2

**Table 3. 3.** The values of the thermal properties of thermo-molded homopolymer films.

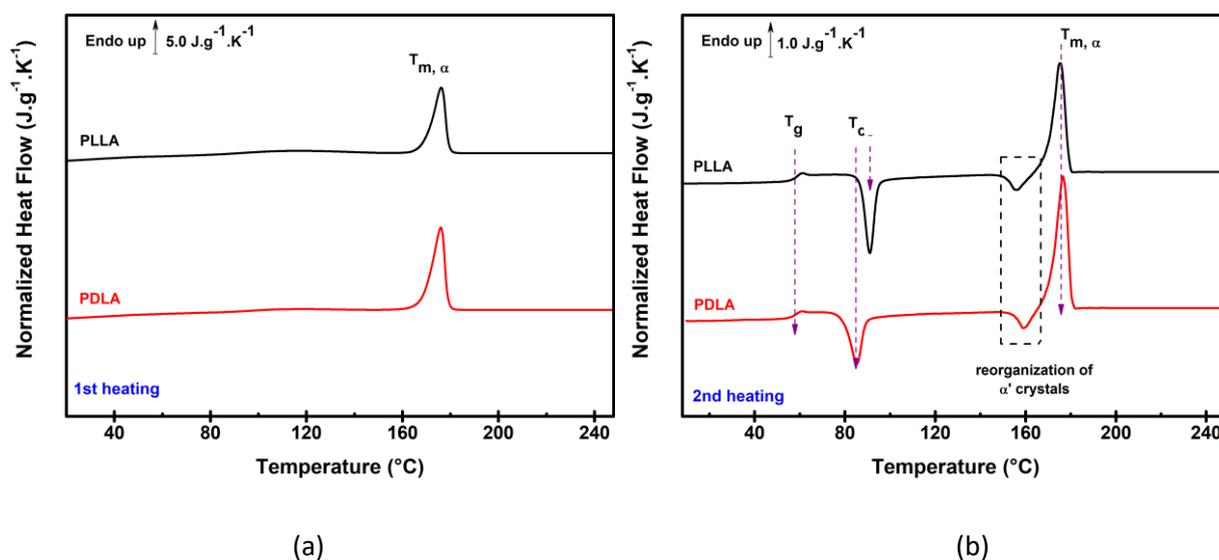
### 3.1.2.2 Semicrystalline homopolymers' film

Semicrystalline homopolymers' films were obtained by solution casting and characterized by DSC, MT-DSC and XRD analysis in order to investigate their thermal properties and microstructure. The DSC curves and thermal characteristics values of homopolymers film are presented in Fig 3.4 and Table 3.4, respectively. The 1<sup>st</sup> heating DSC curves show only endothermic peaks at 176 °C which correspond to the melting peaks of  $\alpha$  homo-crystals of PLLA and PDLA obtained during solvent evaporation.

The obtained homopolymers films are semicrystalline with the  $X_c$  values equal 47 and 41 % for PLLA and PDLA, respectively. In order to study the stability of the homo-crystals 2<sup>nd</sup> heating scan was carried out (Fig 3.4b). Although, there is a  $\alpha'$  crystals formation ( $\sim 85 - 95$  °C), the reformation of  $\alpha$  crystals is confirmed by the melting peak at 176°C. The  $X_c$  values measured during 2<sup>nd</sup> heat are lower than the values obtained at 1<sup>st</sup> heat (Table 3.4). This result

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testifies that the cooling at  $10^{\circ}\text{C}\cdot\text{min}^{-1}$  is not enough to obtain wholly amorphous sample. Therefore, one can say that the  $\alpha$  crystals obtained by solution casting method are rather stable.



**Figure 3. 4.** DSC curves of PLLA and PDLA films obtained by solution casting: (a) 1<sup>st</sup> and (b) 2<sup>nd</sup> heating.

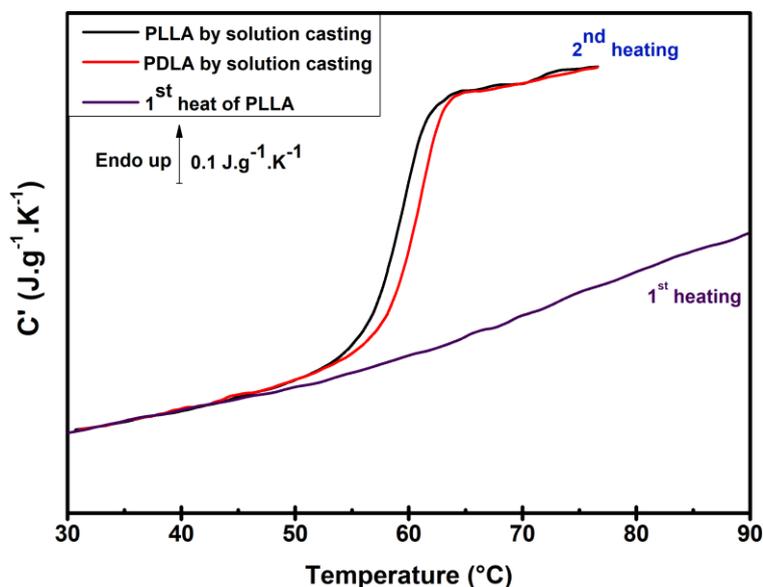
Samples	$\Delta H_m$ (J/g)		$\Delta H_{cc}$ (J/g)		$X_c$ (%)		$T_m$ (°C)	
	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat
PLLA film	44	47	-	26	47	23	176	176
PDLA film	38	40	-	31	41	10	176	176

**Table 3. 4.** The values of the thermal properties of homopolymer films obtained by solution casting.

The properties of the amorphous phase of semicrystalline homopolymers' films were analyzed by MT-DSC to estimate the  $T_g$  and the heat capacity step ( $\Delta C_p$ ). After *in-situ* melting, homopolymer films were quenched at cooling rate  $0.78 \text{ K}\cdot\text{s}^{-1}$  and the MT-DSC curves are presented in Fig 3.5. The  $T_g$  and  $\Delta C_p$  values were determined as explained in section 1.8.1. The values of  $T_g$  and  $\Delta C_p$  shows that homopolymers exhibit the similar glass transition properties as the  $T_g$  values are  $59.3$  and  $60.5$  °C for PLLA and PDLA, respectively, and the  $\Delta C_p$  values are  $0.46 \text{ J}\cdot\text{g}^{-1}\cdot\text{K}^{-1}$  for both homopolymers.

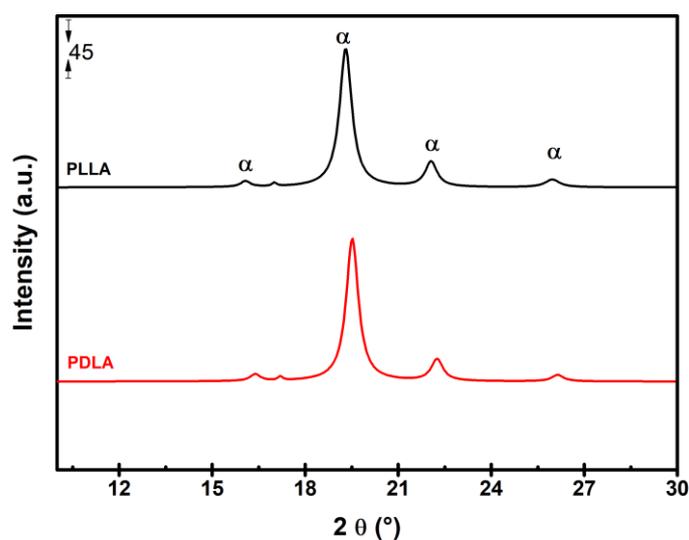
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**Figure 3. 5.** Real component ( $C'$ ) of the complex heat capacity after phase lag correction as a function of temperature for PLLA and PDLA films obtained by solution casting.

The microstructure investigation for homopolymer films obtained by solution casting was carried out by XRD measurements. Homopolymers films have diffractions peaks at  $2\theta = 16.7^\circ, 17.6^\circ, 19.5^\circ, 22^\circ$  and  $26^\circ$  as shown in Fig 3.6. The results of XRD spectra for homopolymer films are in good agreement with the literature as homo-crystals show the reflection at similar  $2\theta$  values [5].



**Figure 3. 6.** XRD patterns for homopolymers films obtained by solution casting.

### 3.2 Stereocomplex PLA

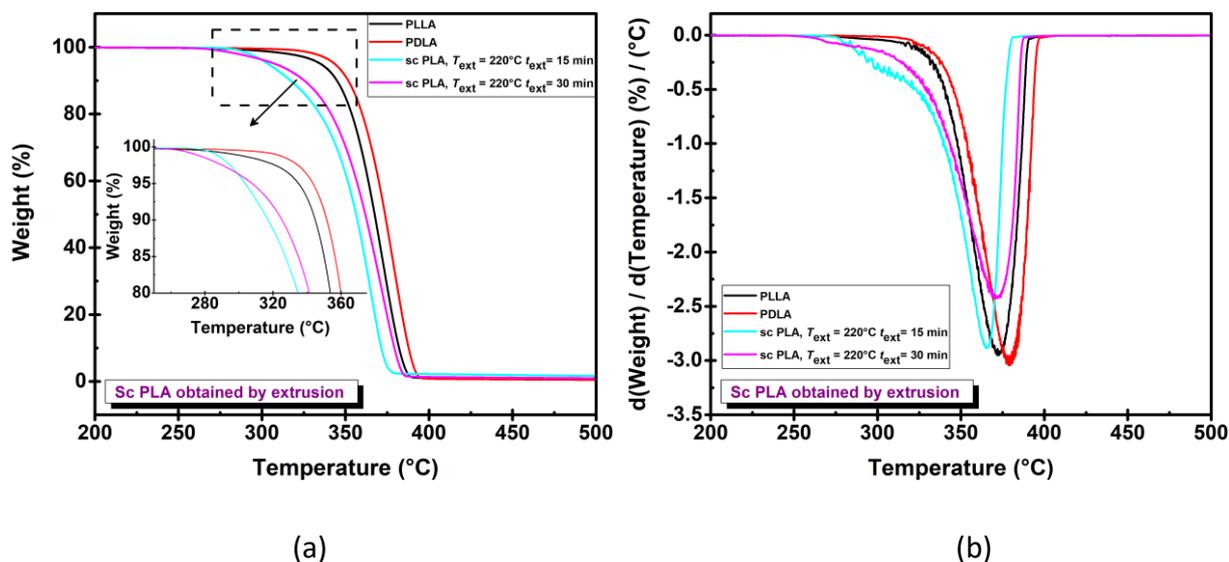
The sc PLA films obtained by extrusion process and solution casting were characterized in order to investigate the influence of the elaboration methods on stereocomplex formation.

#### 3.2.1 Semicrystalline sc PLA by extrusion

Sc PLA samples were analyzed to examine the influence of the extrusion temperature ( $T_{ext}$ ) and extrusion time ( $t_{ext}$ ) on the thermal properties and microstructure of sc PLA.

##### 3.2.1.1 Thermal stability

Thermogravimetric analysis was performed in order to determine the  $T_{deg}$  of sc PLA obtained by extrusion process (Table 3.5). It is shown that the extrusion process has the influence on the  $T_{deg}$  as depicted in Fig 3.7. The degradation of sc PLA obtained by extrusion process is initiated below 300 °C while homopolymers (PLLA and PDLA) start to degrade above 300 °C. In addition, thermal treatment and its time duration decreases the  $T_{deg}$  of sc PLA from 317 °C to 273 °C.



**Figure 3. 7.** TGA curves of sc PLA obtained by extrusion ( $T_{ext} = 220^\circ\text{C}$ ,  $t_{ext} = 15$  and 30 min): (a) thermograms and (b) derivative curves as a function of temperature.

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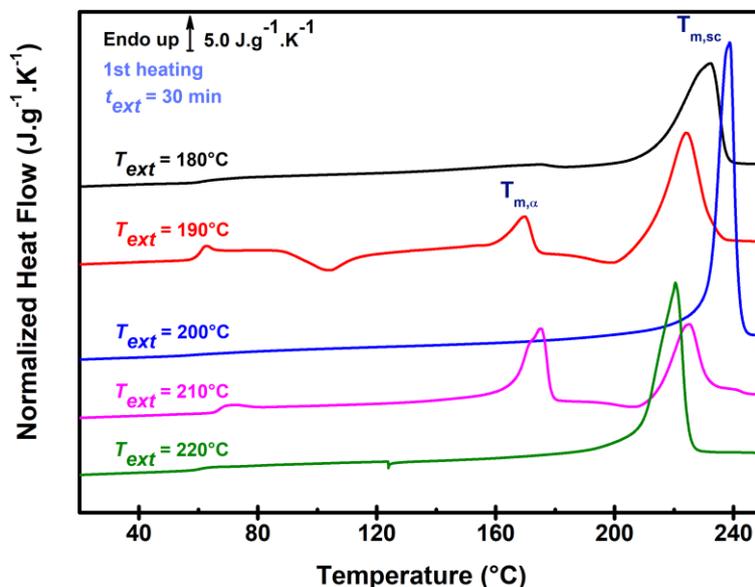
Elaboration conditions	<i>T</i> of 1% weight loss (°C)	<i>T</i> of 50% weight loss (°C)	Maximum of derivative curve (°C)
$T_{ext} = 220\text{ °C}, t_{ext} = 15\text{ min}$	286	356	365
$T_{ext} = 220\text{ °C}, t_{ext} = 30\text{ min}$	273	362	372

**Table 3. 5.** Degradation temperature ( $T_{deg}$ ) of sc PLA obtained by extrusion process.

Such results are supported by the work of Bao et al. [6] on stereocomplex formation by extrusion process. Bao et al. reported that elaboration methods had influence on the thermal degradation temperature of sc PLA as degradation temperature decreased with increasing of extrusion temperature and time. Such diminution of the temperature can be explained by a partial degradation of the polymer chain during the extrusion process.

### 3.2.1.2 Influence of the extrusion temperature ( $T_{ext}$ )

Calorimetric investigations were carried out also for sc PLA obtained by extrusion process in order to study the influence of the extrusion temperature  $T_{ext}$  on the sc PLA formation. As one can see, the increase of the extrusion temperature from 180 °C to 220 °C provokes the formation of sc PLA with different thermal behavior (Fig 3.8). For the blends extruded at  $T_{ext} = 180, 200$  and 220 °C, the pure sc crystals formation was confirmed by the endothermic melting peaks at  $T_{m, sc} = 220 - 238$  °C. In the case of  $T_{ext} = 190$  °C, the cold crystallization of  $\alpha$  crystals  $T_{cc, \alpha} = 103$  °C, the melting of  $\alpha$  crystals  $T_{m, \alpha} = 170$  °C, and the melting of sc crystals  $T_{m, sc} = 224$  °C are observed during the 1<sup>st</sup> heating step. In case of  $T_{ext} = 210$ °C, the melting of  $\alpha$  and sc crystals is observed. The melting enthalpy and the crystallinity degree (Eq 3.1, where the melting enthalpy of 100 % crystalline sc sample is 146 J/g) of extruded samples were determined and presented in Table 3.6. The higher yield of the stereocomplex formation was observed at the extrusion temperature  $T_{ext} = 220$  °C with the crystallinity degree  $X_{c, sc}$  of 55 % and the melting enthalpy sc crystals  $\Delta H_{m, sc}$  of 80 J/g.

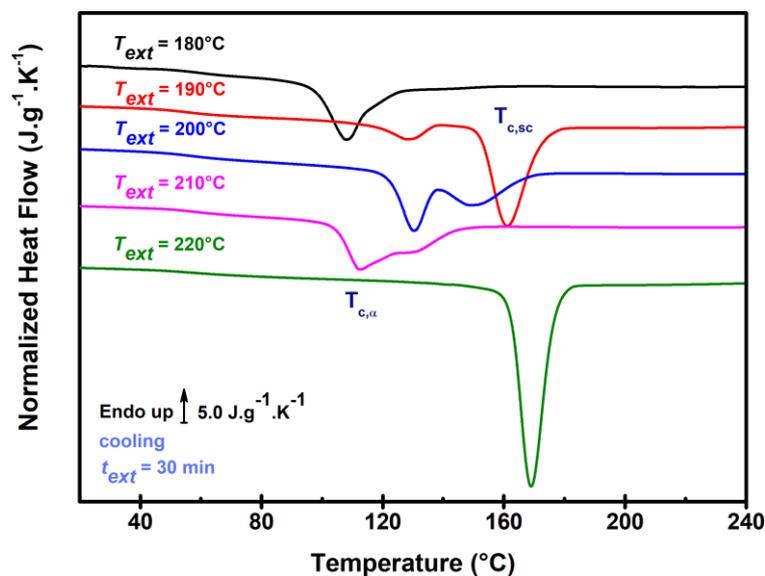


**Figure 3. 8.** The 1<sup>st</sup> heating DSC curves of sc PLA obtained by extrusion process at the extrusion temperature  $T_{ext} = 180 - 220$  °C and the extrusion time  $t_{ext} = 30$  min.

Samples 1 <sup>st</sup> heating	$\Delta H_m$ (J/g)		$X_c$ (%)	
	$\alpha$	sc	$\alpha$	sc
Sc PLA, $T_{ext} = 180$ °C	–	65	–	45
Sc PLA, $T_{ext} = 190$ °C	13	57	14	39
Sc PLA, $T_{ext} = 200$ °C	–	30	–	21
Sc PLA, $T_{ext} = 210$ °C	33	34	35	23
Sc PLA, $T_{ext} = 220$ °C	–	80	–	55

**Table 3. 6.** The values of the melting enthalpy ( $\Delta H_m$ ) and the crystallinity degree ( $X_c$ ) determined from 1<sup>st</sup> heating for sc PLA obtained by extrusion at different extrusion temperature.

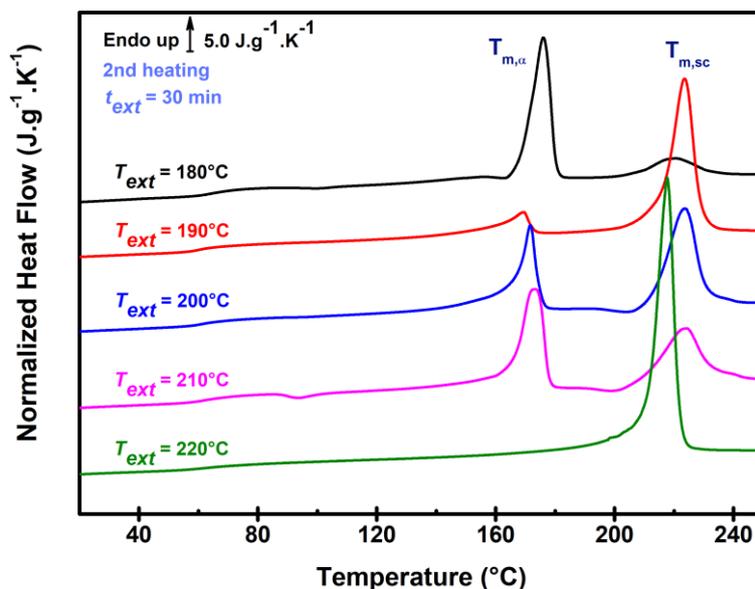
In order to study the crystallization process of  $\alpha$  and sc crystals, the analysis of cooling DSC curves was performed. Fig 3.9 presents the DSC cooling curves at  $|\beta_c| = 10$  K.min<sup>-1</sup> for all samples obtained by extrusion process. In the case of  $T_{ext} = 180$  °C, the crystallization of pure  $\alpha$  crystals was observed at  $T_{c, \alpha} = 107$  °C during the cooling step. For the  $T_{ext} = 190, 200,$  and  $210$  °C, the crystallization of both  $\alpha$  and sc crystals takes place as the exothermic peak is revealed at  $T_{c, \alpha} = 112 - 130$  °C and  $T_{c, sc} = 150 - 162$  °C, respectively (Fig 3.7). In addition, in the case of  $T_{ext} = 220$  °C, the crystallization of sc crystals takes place at  $T_{c, sc} = 169$  °C and no crystallization of  $\alpha$  crystals can be revealed.



**Figure 3. 9.** DSC curves of cooling after 1<sup>st</sup> heating for sc PLA obtained by extrusion at  $T_{ext}$  = 180 – 220 °C,  $t_{ext}$  = 30 min.

During the 2<sup>nd</sup> heating step for  $T_{ext} < 220^{\circ}C$  (presented in Fig 3.10), the formation of  $\alpha$  crystals during cooling is confirmed by the intense melting peak at  $T_{m, \alpha} = 176^{\circ}C$ , whereas the sc crystals melting is observed at  $T_{m, sc} = 220^{\circ}C$ . The decrease in the melting temperature of the sc crystals for  $T_{ext} = 180^{\circ}C$  (i.e. from  $232^{\circ}C$  to  $220^{\circ}C$ ) may be caused by the change in the crystal size reduced by the thermal protocol. It is interesting to note that with the increasing  $T_{ext}$  from  $190^{\circ}C$  to  $210^{\circ}C$ , the formation of  $\alpha$  crystals is increased while stereocomplex formation decreases. This inverse relation can be revealed by the intensity of melting peak of both  $\alpha$  and sc crystals as presented in Table 3.7. These results could be explained by “a low temperature approach” proposed by Bao et al. [6]. The authors reported that blending of the low molecular weight PLLA and PDLA at low temperature ( $T \approx 160 - 170^{\circ}C$ ) could lead to the formation of sc crystals in the blends. On the other hand, Bao et al. also noted that stereocomplexation from high molecular weight polymer ( $MW > 10^5$  kDa) by extrusion should be carried out at high extrusion temperature, i.e. above  $220^{\circ}C$ . However, the higher temperature can cause the thermal degradation of homopolymers during the process, thus a special attention should be paid on the extrusion temperature. In our case, the temperature impact on low molecular weight ( $MW < 10^5$  kDa) PLLA and PDLA homopolymers was observed as the quantity of sc crystals decreased with the increasing temperature. In case of  $T_{ext} = 220^{\circ}C$ ,

the pure sc crystals crystallization during cooling was confirmed by the melting peak at  $T_{m,sc} = 218\text{ }^{\circ}\text{C}$  with the same melting enthalpy and crystallinity degree as during 1<sup>st</sup> heating step (Fig 3.10 and Table 3.7).



**Figure 3. 10.** The 2<sup>nd</sup> heating DSC curves of sc PLA obtained by extrusion process at  $T_{ext} = 180 - 220\text{ }^{\circ}\text{C}$  and  $t_{ext} = 30\text{ min}$ .

Samples 2 <sup>nd</sup> heating	$\Delta H_m$ (J/g)		$X_c$ (%)	
	$\alpha$	sc	$\alpha$	sc
Sc PLA, $T_{ext} = 180\text{ }^{\circ}\text{C}$	37	11	40	8
Sc PLA, $T_{ext} = 190\text{ }^{\circ}\text{C}$	8	55	9	38
Sc PLA, $T_{ext} = 200\text{ }^{\circ}\text{C}$	26	41	28	28
Sc PLA, $T_{ext} = 210\text{ }^{\circ}\text{C}$	36	31	39	22
Sc PLA, $T_{ext} = 220\text{ }^{\circ}\text{C}$	–	80	–	55

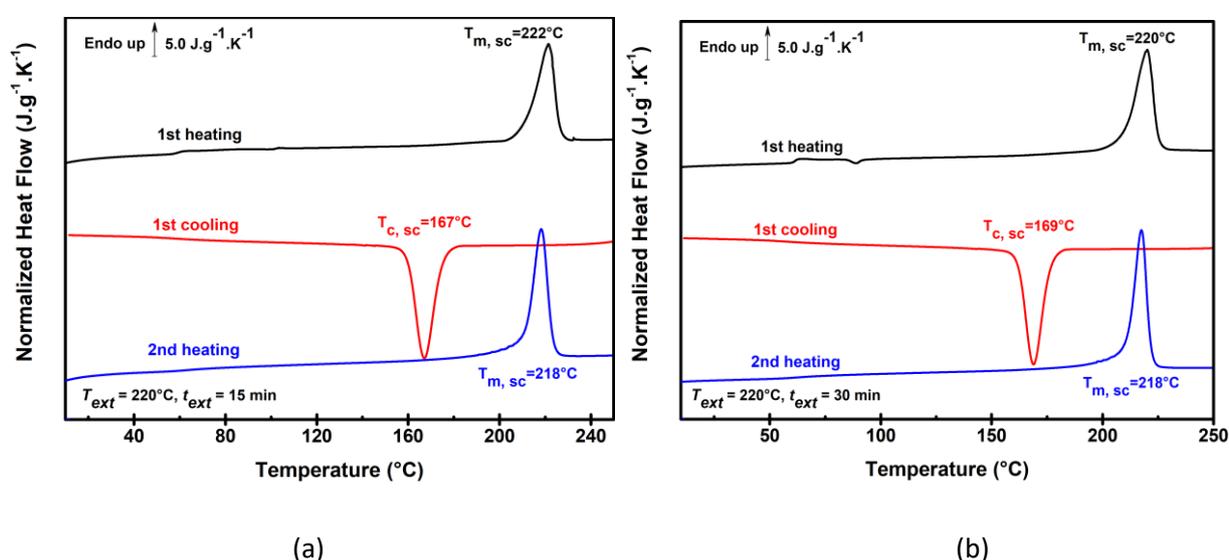
**Table 3. 7.** The values of the melting enthalpy ( $\Delta H_m$ ) and the crystallinity degree ( $X_c$ ) determined from 2<sup>nd</sup> heating for the sc PLA obtained by extrusion at different extrusion temperature.

As one can note from the results in Table 3.7, the highest crystallinity degree of the sc crystals is observed for the sc PLA obtained at  $T_{ext} = 220\text{ }^{\circ}\text{C}$ ,  $t_{ext} = 30\text{ min}$  on the contrary to “a low temperature approach”. Moreover, the only presence of melting of sc crystals at  $T_{m,sc} = 218\text{ }^{\circ}\text{C}$  during the 2<sup>nd</sup> heating step testifies to the highest stability of sc crystals obtained at  $T_{ext} = 220\text{ }^{\circ}\text{C}$  during  $t_{ext} = 30\text{ min}$ . The higher the temperature, the more stable the sc crystals. This

fact can be explained by the nucleation agent exerting distinct effects of stereocomplexation on homocrystallization [7].

### 3.2.1.3 Influence of the extrusion time ( $t_{ext}$ )

In order to study the influence of the extrusion time on the stability of the sc crystals, sc PLA was also elaborated at  $T_{ext} = 220$  °C during  $t_{ext} = 15$  min. As for the sc PLA elaborated during  $t_{ext} = 30$  min, pure sc PLA crystals were obtained during the extrusion process as shown in Fig 3.11.



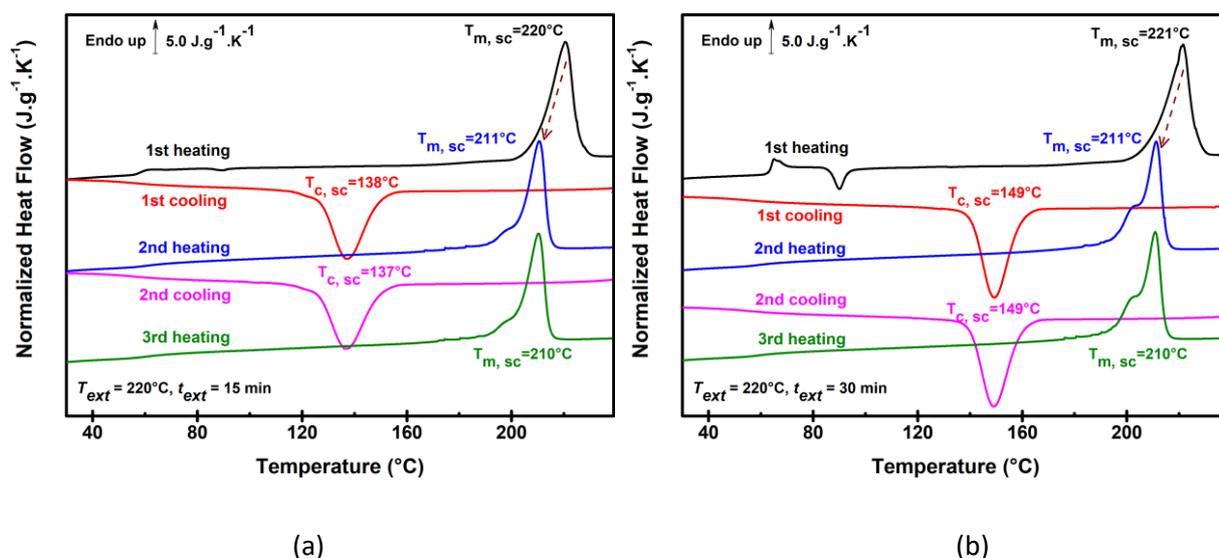
**Figure 3. 11.** DSC curves of sc PLA obtained by extrusion process at  $T_{ext} = 220$  °C during the extrusion time (a)  $t_{ext} = 15$  min and (b)  $t_{ext} = 30$  min.

No formation of  $\alpha$  crystals is observed during the cooling as the curve of the 2<sup>nd</sup> heating step is characterized by only melting peak of sc crystals at  $T_{m,sc} = 218$  °C for both studied extrusion time. The sc PLA elaborated at  $t_{ext} = 15$  min has the same values of melting enthalpy ( $\Delta H_m = 80$  J/g) and the crystallinity degree ( $X_c = 55\%$ ) as the one elaborated at  $t_{ext} = 30$  min. Therefore, one can conclude that the optimum extrusion conditions of pure sc crystals formation are  $T_{ext} = 220$  °C and  $t_{ext} = 15$  min.

### 3.2.1.4 Stability of sc crystals obtained by extrusion

Isothermal step during  $t_{iso} = 120$  min at  $T_{iso} = 260$  °C was performed between the 1<sup>st</sup> heating and 1<sup>st</sup> cooling step. After this isothermal step, the samples were cooled down to 0 °C

and heated one more time to 260 °C (10 K.min<sup>-1</sup>). As shown in Fig 3.10, sc crystals in the sc PLA are thermally stable. In sc PLA extruded during 15 and 30 min, the presence of pure sc PLA is confirmed by the melting peak at  $T_{m,sc} = 221$  °C during 1<sup>st</sup> heating step. During the cooling step, the sc crystal crystallization is observed at  $T_{m,sc} = 137$  °C and 149 °C for the blend extruded at 220 °C during 15 min and 30 min, respectively, (Fig 3.12a and 3.12b, respectively).



**Figure 3. 12.** DSC curves of sc PLA obtained by extrusion process at  $T_{ext} = 220$  °C during (a)  $t_{ext} = 15$  min and (b)  $t_{ext} = 30$  min. Isothermal step was applied at  $T_{iso} = 260$  °C during 120 min between the 1<sup>st</sup> heating step and the 1<sup>st</sup> cooling step.

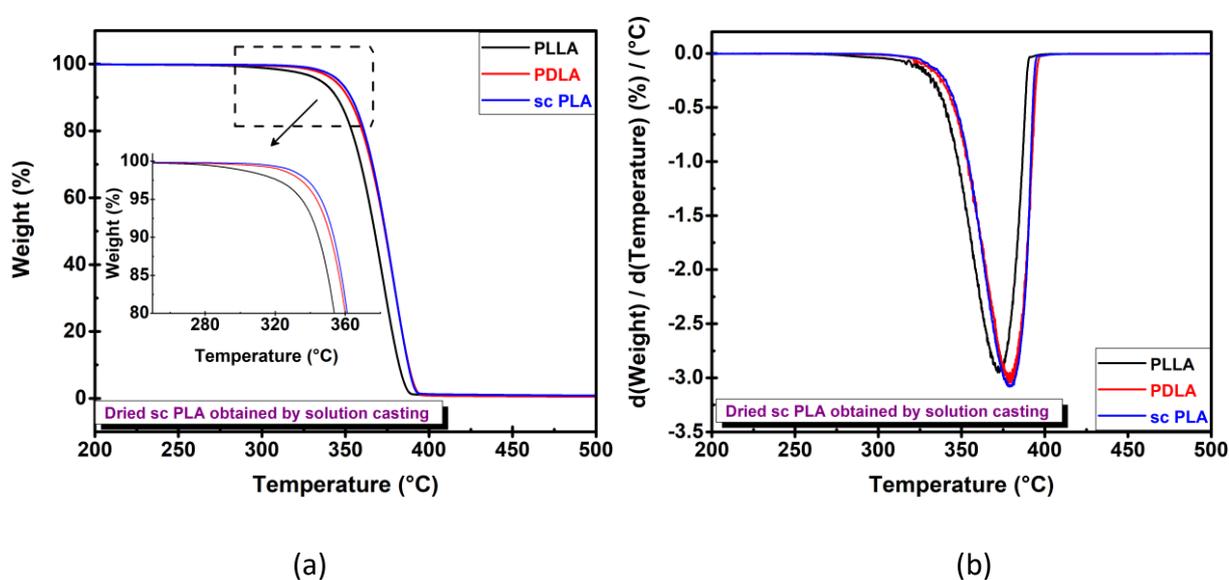
The heating and cooling cycles were carried out for several times in order to confirm the thermal stability and thermal characteristics of sc PLA. The decrease of 10 °C in the melting temperature can be explained by the change in the crystal size of sc crystals during the heat-cool processes.

### 3.2.2 Semicrystalline sc PLA by solution casting

Sc PLA obtained by solution casting was analyzed to investigate the influence of the polymer concentration and the ratio of the homopolymers (PLLA and PDLA) on the sc PLA formation.

### 3.2.2.1 Thermal stability

TGA was performed to determine the  $T_{deg}$  in order to examine the influence of solution casting method on thermal degradation. The solution casting method has no significant influence on the  $T_{deg}$  as one can see from Fig 3.13 and Table 3.8. Sc PLA obtained by solution casting starts to degrade above 300 °C as well as homopolymers (PLLA and PDLA), while the degradation of sc PLA obtained by extrusion process is initiated below 300 °C (section 3.2.1.1). Thus, the thermal degradation of sc PLA depends strongly on the elaboration method used.



**Figure 3. 13.** TGA curves of pure PLLA and PDLA pellets and sc PLA obtained by solution casting: (a) thermograms and (b) derivative curves as a function of temperature.

Elaboration conditions	$T$ of 1% weight loss (°C)	$T$ of 50% weight loss (°C)	Maximum of derivative curve (°C)
PLLA pellet	296	367	373
PDLA pellet	321	373	378
PLLA:PDLA = 50:50, $d = 3$ g/dL	317	362	379

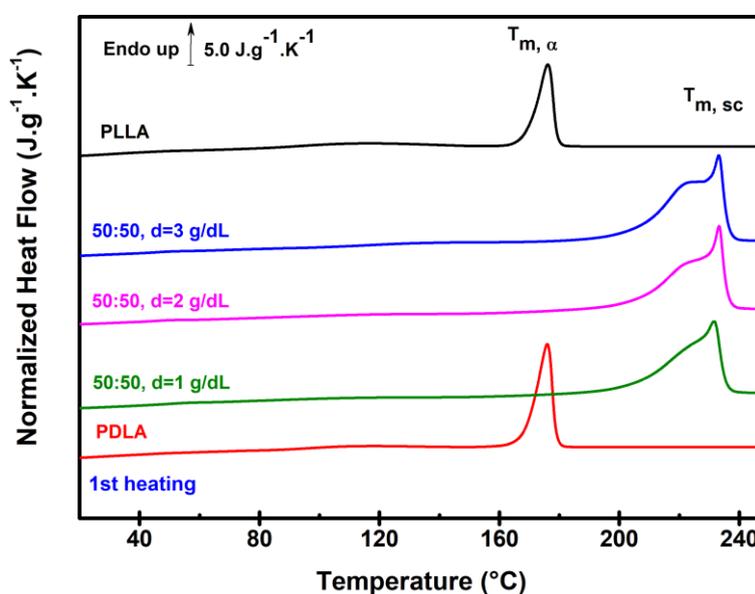
**Table 3. 8.** Degradation temperature ( $T_{deg}$ ) of sc PLA (PLLA:PDLA = 50:50) obtained by solution casting.

### 3.2.2.2 Influence of the polymer concentration

The influence of polymer concentration was investigated by DSC and XRD analysis to define the optimum conditions for the sc crystals elaboration. During DSC analysis the

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stereocomplexation is characterized by means of the melting peaks of sc,  $\alpha$  and  $\alpha'$  crystals. Fig 3.14 presents the normalized heat flow curves of sc PLA films obtained by solution casting at 50:50 ratio of PLLA:PDLA with the concentration of 1, 2 and 3 g/dL. The curves of pure homopolymers (PLLA and PDLA) are given for comparison. The endothermic peaks at temperature of 176 °C and 233 °C correspond to the melting of  $\alpha$  crystals and sc crystals, respectively. It should be noted that the shape of sc PLA melting peaks show different behavior compared to homopolymers melting peaks. The doubled melting peak testifies to the not homogenous distribution of the crystals size in sc PLA samples.



**Figure 3. 14.** DSC curves of sc PLA film obtained by solution casting (PLLA:PDLA = 50:50) as a function of polymer concentration.

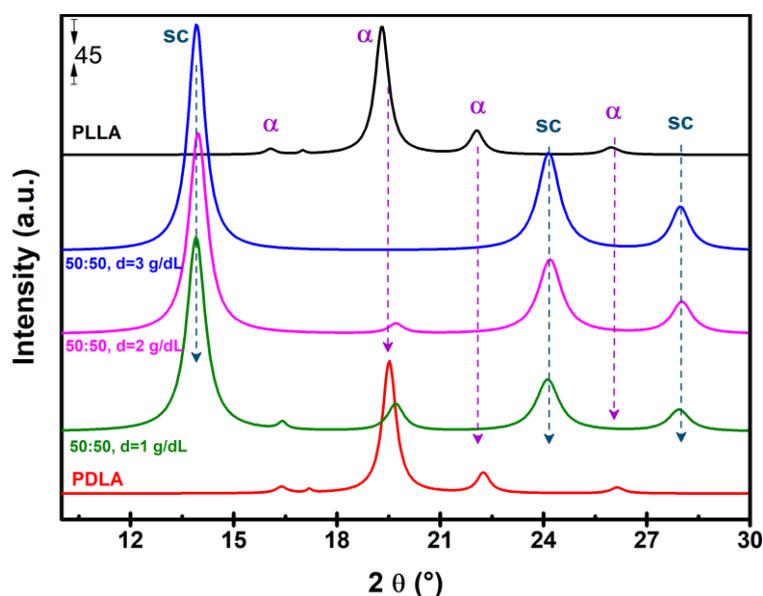
Table 3.9 presents the crystallinity degree values. In the sc PLA film at 50:50 ratio of PLLA:PDLA, the pure sc crystallites were obtained whatever the polymer concentration. As shown in Fig 3.14 and Table 3.9, the polymer concentration has no significant influence on the stereocomplexation.

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Samples	$\Delta H_m$ (J/g)		$X_c$ (%)	
	$\alpha$	sc	$\alpha$	sc
PLLA	44	–	47	–
PLLA:PDLA = 50:50, d = 1 g/dL	–	75	–	53
PLLA:PDLA = 50:50, d = 2 g/dL	–	80	–	56
PLLA:PDLA = 50:50, d = 3 g/dL	–	77	–	54
PDLA	38	–	41	–

**Table 3. 9.** The values of the melting enthalpy ( $\Delta H_m$ ) and the crystallinity degree ( $X_c$ ) as a function of polymer concentration.

In addition, the microstructure of sc PLA films obtained at the ratio of 50:50 PLLA:PDLA with the concentration of 1, 2 and 3 g/dL was investigated and XRD spectra are presented in Fig 3.15. Sc PLA shows diffraction peaks at  $2\theta = 14^\circ$ ,  $24^\circ$  and  $28^\circ$  while homopolymers PLLA and PDLA have peaks at  $2\theta = 19.5^\circ$ ,  $22^\circ$  and  $26^\circ$ .



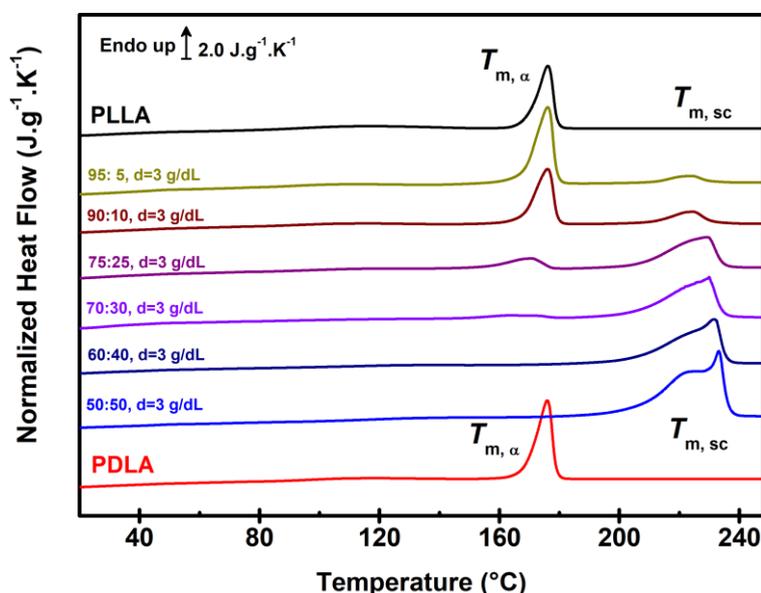
**Figure 3. 15.** XRD patterns of the sc PLA film obtained by solution casting at the ratio of 50:50 PLLA:PDLA with the concentration of 1, 2, and 3 g/dL.

Although, the  $\alpha$  crystals presence in the film of 50:50 ratio of PLLA:PDLA with the concentration of 1 and 2 g/dL was not confirmed by DSC measurements, the XRD spectra of these samples show the  $\alpha$  crystal existence confirmed by small intensive peaks at  $17^\circ$  and  $19^\circ$ .

Therefore, one may noticed that the polymer concentration has slight influence on the homocrystals presence in the obtained film of sc PLA.

### 3.2.2.3 Influence of the ratio of homopolymers

In order to study the influence of the homopolymers' ratio on the stereocomplexation reaction, the films with different ratio of homopolymers (PLLA:PDLA) were elaborated. The results of 1<sup>st</sup> heating scan are presented in Fig 3.16. The presence of  $\alpha$  crystals was confirmed by the endothermic melting peak at  $T_{m, \alpha} = 176$  °C. At the same time, the peak at  $T_{m, sc} = 225 - 235$  °C testifies to the sc crystals presence. Thermal analysis results show that the homocrystallization occurs simultaneously due to the unpaired homopolymers chain when deviating from equimolar ratio of PLLA and PDLA. According to the obtained stereocomplex crystalline structure, PLLA and PDLA chains are paired in each units with the equilateral triangles structure. Therefore, equimolar blending ratio (50:50) is found to be the optimum condition for stereocomplex crystals formation. The result of pure stereocomplex formation at 50:50 ratio of homopolymers was also reported by Ikada et al. [8].



**Figure 3. 16.** DSC curves of sc PLA film obtained by solution casting at constant polymer concentration ( $d= 3$  g/dL) as a function of homopolymer ratio.

The crystallinity degree of  $\alpha$  and sc crystals was calculated by Eq 3.1 and the obtained values are gathered in Table 3.10. As shown in Fig 3.16 and Table 3.10, the ratio of

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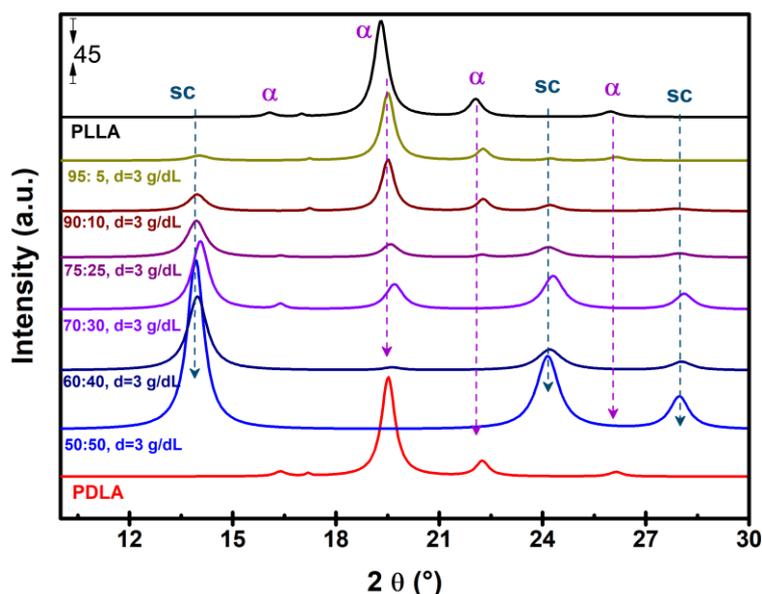
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homopolymers has significant influence on the stereocomplex formation. The decrease in the melting enthalpy and the crystallinity degree of sc crystals was observed with the decreasing of the PDLA content in the racemic mixture of PLLA and PDLA. Tsuji et al. [9-11] indicated that the melting peaks of both homo crystals and sc crystals were observed when PDLA fraction ( $X_D$ ) was between 0.1 to 0.3 and 0.7 to 0.9, while the pure stereocomplex was detected for  $X_D$  between 0.4 and 0.6 with maximum of the melting enthalpy ( $\Delta H_m$ ) at  $X_D = 0.5$ . The pure homocrystallization occurred near  $X_D = 0$  and 1. In our case, the optimum blending ratio of homopolymers for pure stereocomplex formation was found to be 60:40 and 50:50 (Table 3.10).

Samples	$\Delta H_m$ (J/g)		$X_c$ (%)	
	$\alpha$	sc	$\alpha$	sc
PLLA	44	–	47	–
PLLA:PDLA = 95:05, d = 3 g/dL	42	7	45	5
PLLA:PDLA = 90:10, d = 3 g/dL	33	13	35	9
PLLA:PDLA = 75:25, d = 3 g/dL	11	41	12	29
PLLA:PDLA = 70:30, d = 3 g/dL	4	48	4	34
PLLA:PDLA = 60:40, d = 3 g/dL	–	64	–	45
PLLA:PDLA = 50:50, d = 3 g/dL	–	75	–	53
PDLA	38	–	41	–

**Table 3. 10.** The melting enthalpy ( $\Delta H_m$ ) and crystallinity degree ( $X_c$ ) values as a function of homopolymer ratio determined from 1<sup>st</sup> heating scan.

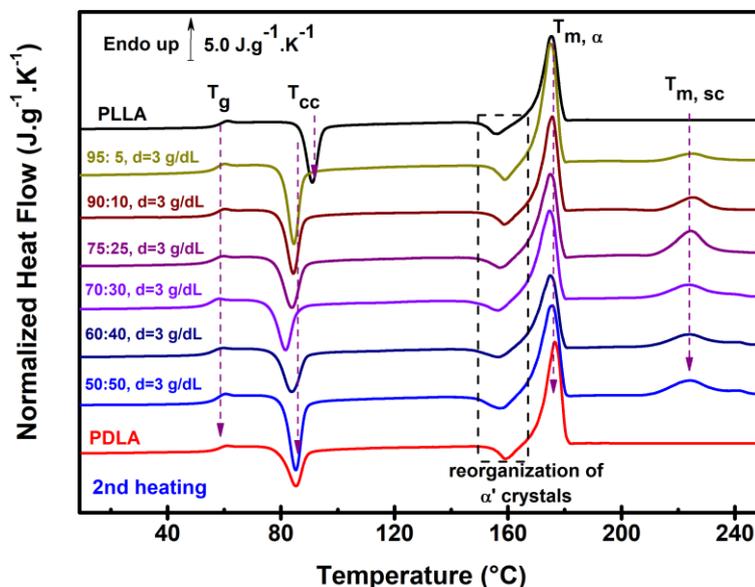
The XRD patterns of films with different ratio of PLLA:PDLA (i.e. from 50:50 up to 95:5) are presented in Fig 3.17. As one can see, when deviating from the equivalent ratio of homopolymers, the formation of homocrystals is observed. The existence of sc crystals and  $\alpha$  crystals is revealed by the reflection peaks of  $\alpha$  crystals ( $2\theta = 19.5^\circ, 22^\circ$  and  $26^\circ$ ) and sc crystals ( $2\theta = 14^\circ, 24^\circ$  and  $28^\circ$ ). The results of XRD spectra are in good accordance with the literature, as the  $\alpha$  crystals show the reflection at  $2\theta = 16.6^\circ, 19.1^\circ$  and  $17^\circ, 19^\circ$  [5] and the sc crystals show the reflection at  $2\theta = 12^\circ, 21^\circ, 24^\circ$  [5] and  $12^\circ, 20.9^\circ, 24^\circ$  [12].



**Figure 3. 17.** XRD patterns of the PLLA/PDLA film obtained by solution casting at the different ratio of PLLA:PDLA.

The stability of the sc PLA was investigated during 2<sup>nd</sup> heating scan of DSC measurements (Fig 3.18). It was found that the sc PLA crystals obtained by solution casting were not stable, as already during the 2<sup>nd</sup> heating up to 250 °C the crystallization and melting of  $\alpha$  crystals was visible and confirmed by cold-crystallization and melting peak. Moreover, the intensity of melting peak  $T_{m, sc}$  at 223 °C, which corresponds to the sc crystals melting, decreases significantly after heating (Fig 3.18). The values of the melting enthalpy and crystallinity degree drastically reduce for sc crystals. The crystallinity degree of sc crystals decreases from 54 % to 10 % even for the sample with 50:50 and 60:40 ratio of PLLA:PDLA during the 2<sup>nd</sup> heating scan, as shown in Table 3.11. In addition, the  $\alpha'$  crystals formation is observed with cold-crystallization at  $\approx 85$  °C (the cold-crystallization at temperature  $T < 100$  °C leads to the formation of  $\alpha'$  crystals form consisting in disordered crystals with hexagonal packing [1, 2]). The reorganization of  $\alpha'$  imperfect crystals into  $\alpha$  form crystals was observed by the exothermic peak before melting at temperature  $T \approx 160$  °C. The presence of sc PLA crystals in PLLA/PDLA films was confirmed by  $T_m$  value found between 223 °C and 233 °C, depending on the homopolymers' ratio (Fig 3.16 and 3.18).

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**Figure 3. 18.** The 2<sup>nd</sup> heating DSC curves of sc PLA film obtained by solution casting at different ratio of PLLA:PDLA.

The DSC results confirm that the heating of the blends leads to homopolymer crystallization whatever the ratio of homopolymers. Only under certain experimental conditions the pure sc PLA crystals were obtained (i.e. without  $\alpha$  and  $\alpha'$  crystals) by solution casting, namely at 50:50 and 60:40 ratio of PLLA:PDLA. Moreover, it can be pointed out that sc material is not stable, as the  $\alpha'$  crystals could appear during cooling and 2<sup>nd</sup> heating (after 1<sup>st</sup> melting, Fig 3.18).

Samples 2 <sup>nd</sup> heating	$\Delta H_{cc}$ (J/g)		$\Delta H_m$ (J/g)		$X_c$ (%)	
	$\alpha$	sc	$\alpha$	sc	$\alpha$	sc
PLLA	26	–	47	–	23	–
PLLA:PDLA = 60:40, d = 3 g/dL	28	–	39	14	12	10
PLLA:PDLA = 50:50, d = 1 g/dL	31	–	37	16	6	11
PLLA:PDLA = 50:50, d = 2 g/dL	31	–	37	14	6	10
PLLA:PDLA = 50:50, d = 3 g/dL	31	–	37	15	6	11
PDLA	31	–	40	–	10	–

**Table 3. 11.** The melting enthalpy ( $\Delta H_m$ ) and the crystallinity degree ( $X_c$ ) values determined from 2<sup>nd</sup> heating scan as a function of PLLA:PDLA ratio.

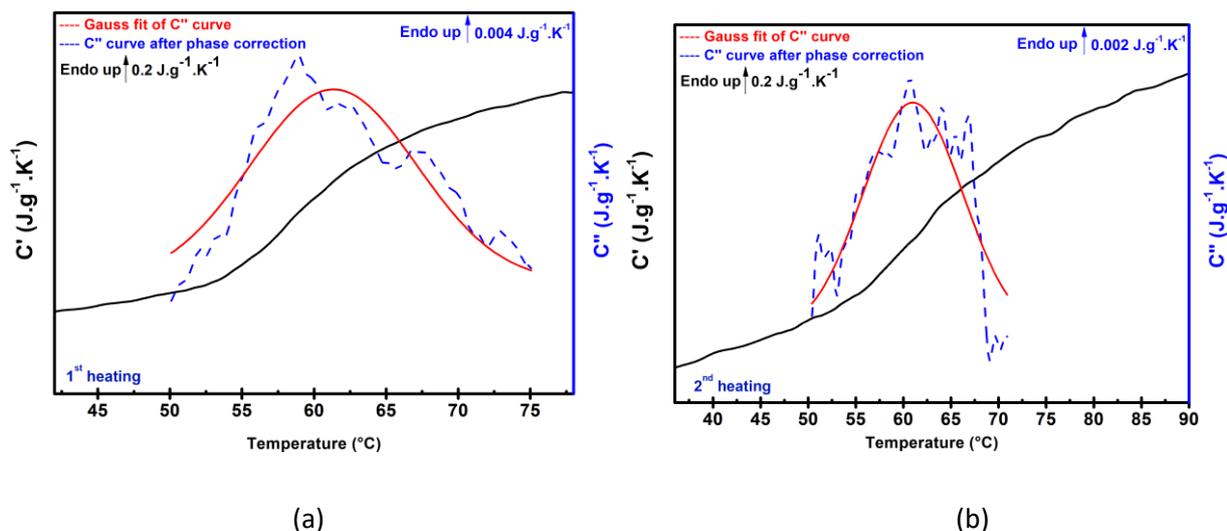
### **3.3 Amorphous phase of stereocomplex PLA**

MT-DSC analysis was performed to investigate the amorphous character and to determine the calorimetric glass transition temperature of the samples elaborated by solution casting and extrusion process. Since the material barrier properties are linked with the molecular mobility and free volume, the Donth's approach [13] was applied in order to investigate the free volume and the CRR size (as detailed in section 1.8.2). MT-DSC measurements were carried out in heat-cool mode, and obtained curves showed the real ( $C'$ ) and the imaginary ( $C''$ ) parts of the complex heat capacity as a function of temperature (detailed in section 2.2.3)

#### **3.3.1 MT-DSC analysis of sc PLA obtained by extrusion process**

It was found by the conventional DSC analysis that the optimum conditions of the stereocomplex formation by extrusion process were the extrusion temperature  $T_{ext} = 220\text{ }^{\circ}\text{C}$  and the extrusion time  $t_{ext} = 15\text{ min}$ . Therefore, MT-DSC measurements were carried out for this sc PLA blend. MT-DSC curves of sc PLA are presented in Fig 3.19. The values of glass transition temperature estimated from the 1<sup>st</sup> and 2<sup>nd</sup> heating step of MT-DSC measurements are presented in Table 3.12. The CRR size of semicrystalline sc PLA films is estimated by Eq 1.20 to be 1.6 nm and 1.3 nm (from 1<sup>st</sup> heating and 2<sup>nd</sup> heating), respectively, and is discussed later. Sc PLA obtained by extrusion process has similar glass transition temperature  $T_g$  (60 – 62  $^{\circ}\text{C}$ ) as amorphous neat PLA and homopolymers, but it has different heat capacity step: 0.24  $\text{J}\cdot\text{g}^{-1}\cdot\text{K}^{-1}$  compared to 0.15  $\text{J}\cdot\text{g}^{-1}\cdot\text{K}^{-1}$  for homopolymers. The results show that the amorphous phase of sc PLA is not interlocked by the sc crystallites during the stereocomplexation by extrusion process. The DSC curves reveal that sc PLA obtained by extrusion process is not quenched to obtained wholly amorphous sc PLA by the cooling rate of  $|\beta_c| = 0.78\text{ K}\cdot\text{min}^{-1}$  which is the maximum cooling rate for the used apparatus (Q100 TA Instruments). Therefore, sc PLA sample is semicrystalline during the 2<sup>nd</sup> heating step. Therefore, the minimum required cooling rate for quenching of sc PLA obtained by extrusion was investigated by FSC. It was found that sc PLA obtained by extrusion process should be vitrified at the scanning rate  $|\beta_c|$  higher than 200  $\text{K}\cdot\text{min}^{-1}$ .

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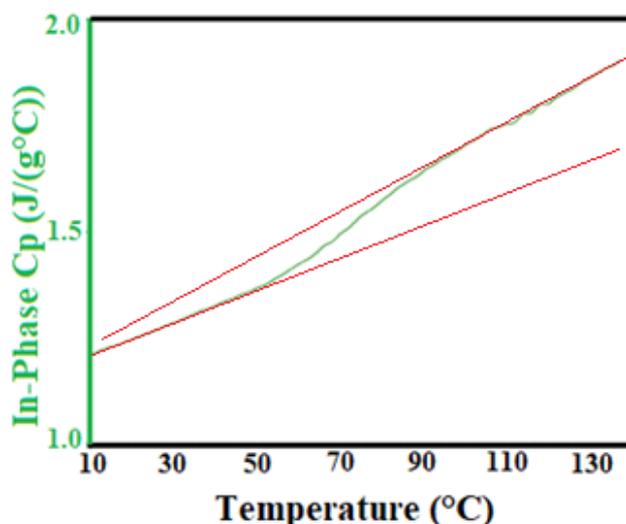
**Figure 3. 19.**  $C'$  and  $C''$  curves of sc PLA obtained by extrusion: (a) the 1<sup>st</sup> heating and (b) 2<sup>nd</sup> heating scan.

PLLA/PDLA blend by extrusion	$T_g$ (°C)	$\Delta C_p$ ( $J.g^{-1}.K^{-1}$ )	$T_{max}$ of Gaussian fit	$\xi$ (nm)	$X_c$ (%)
1 <sup>st</sup> heating step	60	0.24	61	1.6	55
2 <sup>nd</sup> heating step	62	0.15	61	1.3	55

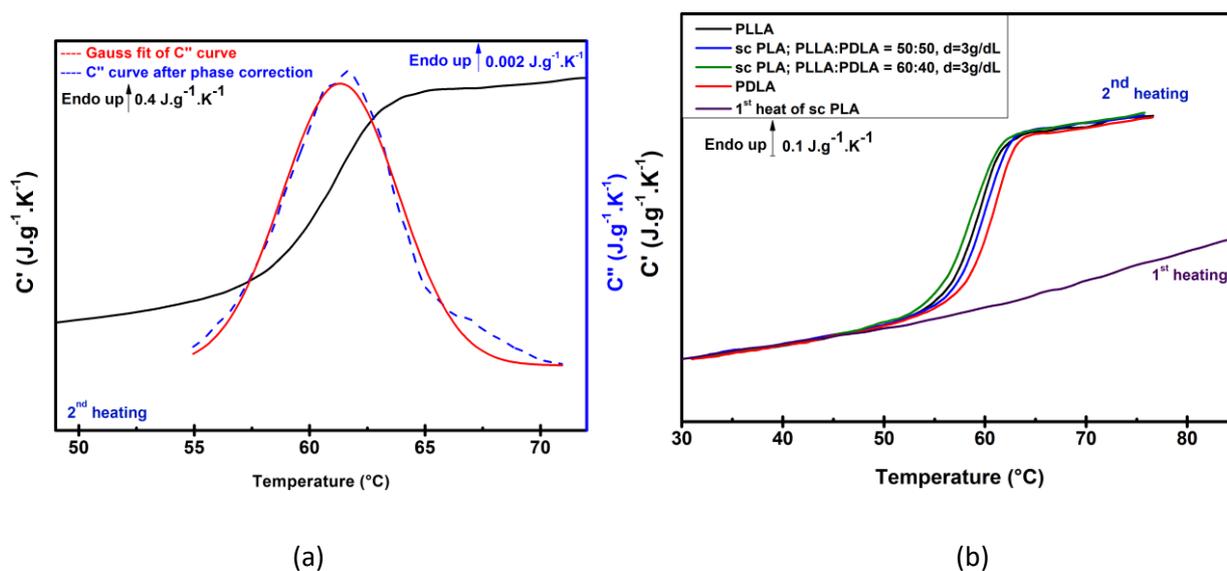
**Table 3.12.** Glass transition parameters of sc PLA obtained by extrusion process.

### 3.3.2 MT-DSC analysis of sc PLA obtained by solution casting

Since the presence of pure sc crystals was observed in PLLA/PDLA film at the ratio of 50:50 and 60:40 of PLLA:PDLA, the glass transition investigations were carried out for these samples. The 1<sup>st</sup> heating ramp of MT-DSC measurements of sc PLA film obtained by solution casting exhibits low heat capacity step due to high  $X_c$  (Fig 3.20). Therefore, only the 2<sup>nd</sup> heating step after quenching at  $|\beta_c| = 0.78 \text{ K.min}^{-1}$  will be presented. Fig 3.21a presents the real ( $C'$ ) and the imaginary ( $C''$ ) parts of complex heat capacity of sc PLA film (PLLA:PDLA = 50:50). Fig 3.21b reveals the glass transition and the heat capacity step of amorphous sc PLA film with PLLA:PDLA ratio of 50:50 and 60:40. The determined values of the glass transition and the CRR size are presented in Table 3.13. The CRR size of semicrystalline (from 1<sup>st</sup> heating) and fully amorphous (from 2<sup>nd</sup> heating) sc PLA films was estimated by Eq 1.20 to be 0.8 nm and 2.9 nm, respectively. The number of structural units in CRR ( $N_\alpha$ ) was determined for the semicrystalline and totally amorphous PLLA/PDLA films (Table 3.13) according to Eq 1.21.



**Figure 3. 20.**  $C'$  curve of semicrystalline sc PLA film (PLLA:PDLA = 50:50) obtained by solution casting during the 1<sup>st</sup> heating scan of MT-DSC measurement.



**Figure 3. 21.** MT-DSC curves of sc PLA obtained by solution casting: (a) the real ( $C'$ ) and the imaginary ( $C''$ ) parts of the complex heat capacity as a function of temperature, (b) the glass transition of the amorphous sc PLA films.

Wholly amorphous sc PLA has the similar heat capacity step as amorphous neat PLA [14], PLLA and PDLA. Wholly amorphous sc PLA has the glass transition temperature  $T_g$  at 61 °C and the heat capacity step of  $0.54 \text{ J.g}^{-1}.\text{K}^{-1}$ , whereas neat PLA and homopolymers (PLLA, PDLA) have

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the glass transition temperature  $T_g$  at 60 °C, the heat capacity step of neat PLA and homopolymers are 0.51 J.g<sup>-1</sup>.K<sup>-1</sup> and 0.46 J.g<sup>-1</sup>.K<sup>-1</sup>, respectively. It should be noted that the glass transition of semicrystalline sc PLA film obtained by solution casting is rather wide and the temperature transition range ( $\delta T > 10$  °C) is out of the limits of Donth's approach (shown in Fig 3.20), thus the  $N_\alpha$  value of semicrystalline sc PLA film is too small ( $N_\alpha = 5$ ).

PLLA/PDLA film by solution casting	$T_g$ (°C)	$\Delta C_p$ (J.g <sup>-1</sup> .K <sup>-1</sup> )	$T_{max}$ of Gaussian fit (°C)	$\delta T$	$\xi$ (nm)	$N_\alpha$
semicrystalline	69	0.12	58	16.9	0.8	5
amorphous	61	0.54	61	2.5	2.9	262

**Table 3. 13.** Glass transition parameters of sc PLA obtained by solution casting method determined from MT-DSC measurements.

### 3.3.3 Comparing the amorphous phase species of sc PLA by “A 3-phase model”

During the crystallization of polymer, the amorphous region volume is regularly diminished and the decoupling between crystalline and amorphous phases remains generally incomplete due to the macromolecular length [15]. Such incomplete coupling leads to decrease in chain mobility of amorphous phase. It is suggested to describe the semicrystalline polymers by a 3-phase model including the crystalline phase, the mobile amorphous phase (MAP) and the rigid amorphous fraction (RAF) [16, 17]. According to Androsch and Wunderlich [18], the RAF amount corresponds to a number of the coupling between the crystalline and the amorphous phase. Whereas the MAP remains inside the lamella stack, the RAF is obtained as an interfacial nanolayer between the lamellae and MAP [19]. The in-phase components  $C'$  are obtained and depicted in Fig 3.19 and 3.20 for the semicrystalline sc PLA obtained by extrusion and solution casting, respectively. Typically, the MAP degree  $X_{MAP}$  from the  $\Delta C_p$  step at  $T_g$  can be calculated as [14]:

$$X_{MAP} = \frac{\Delta C_p}{\Delta C_p^0} \quad (3.2)$$

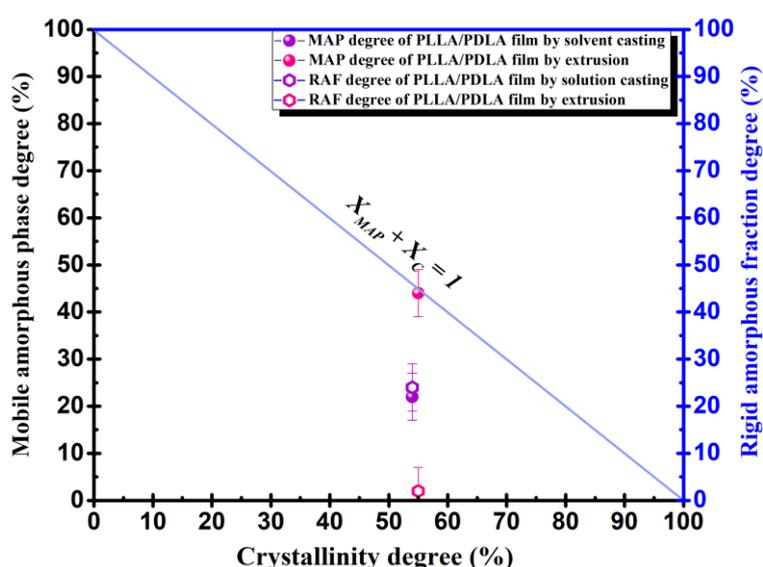
where  $\Delta C_p = C'_{liquid} - C'_{solid}$  is the thermal heat capacity step at  $T_g$  for semicrystalline sc PLA samples and  $\Delta C_p^0$  is that of the wholly amorphous sample. As shown in Table 3.14, and Fig. 3.22, the total of the  $X_{MAP}$  and  $X_c$  is far from the equation line  $X_{MAP} + X_c = 1$ . Therefore, the RAF

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must be taken into account in a 3-phase model. In this case  $X_{MAP} + X_c + X_{RAF} = 1$ , where  $X_{RAF}$  corresponds to the RAF content.

Sc PLA samples	$\Delta C_p$ ( $J \cdot g^{-1} \cdot K^{-1}$ )		$X_{MAP}$ (%)		$X_{RAF}$ (%)		$X_c$ (%)	
	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat	1 <sup>st</sup> heat	2 <sup>nd</sup> heat
Solution casting	0.12	0.54	22	100	24	-	54	-
Extrusion	0.24	0.15	44	28	1	17	55	55

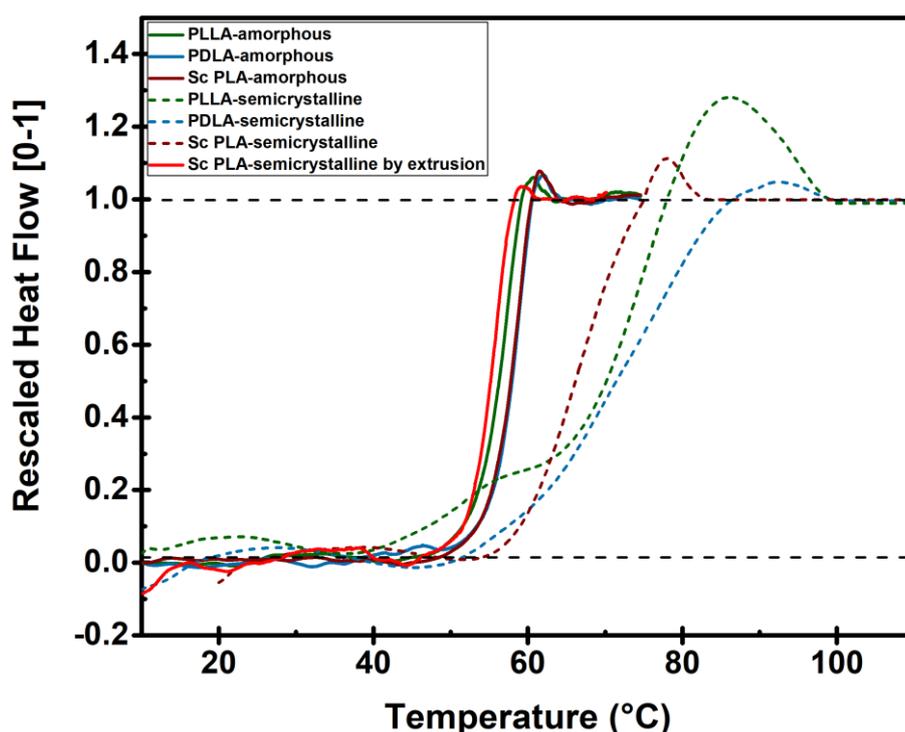
**Table 3. 14.** The values of the degree of mobile amorphous phase  $X_{MAP}$ , rigid amorphous fraction  $X_{RAF}$ , and the crystallinity  $X_c$  of sc PLA obtained by solution casting and extrusion.



**Figure 3. 22.** The evolution of the degree of mobile amorphous phase ( $X_{MAP}$ ) and rigid amorphous fraction ( $X_{RAF}$ ) as a function of crystallinity degree ( $X_c$ ) of sc PLA obtained by solution casting (purple symbols) and by extrusion (pink symbols).

On the other hand, such results show that the mobile amorphous phase of sc PLA obtained by extrusion is higher than that obtained by solution casting, whereas the rigid amorphous phase of sc PLA obtained by extrusion is negligible and lower than the one of sc PLA obtained by solution casting. In order to examine the relaxation of amorphous phase of semicrystalline sc PLA obtained by both solution casting and extrusion, the heat capacity step was normalized to be in 0 to 1 scale (Fig 3.23). As seen in Fig 3.23, the amorphous phase of sc PLA obtained by extrusion ( $T_{ext} = 220^\circ C$ ,  $t_{ext} = 30$  min) relaxes independently from the existence of stereocomplex crystalline structure, while the amorphous phase of sc PLA obtained by

solution casting relaxes at higher temperature due to crystalline structure hindrance. It is found that the mechanical and thermal impact during the extrusion process lead to a better decoupling between the crystalline and the amorphous phase. Such difference can be explained by the crystallization process which is obtained from the molten state during the extrusion when is occurred at room temperature during the solution casting method.



**Figure 3. 23.** Rescaled heat flow curves of amorphous and semicrystalline homopolymers and sc PLA obtained by solution casting and extrusion process.

### Conclusion

Stereocomplex PLAs were successfully elaborated by both solution casting and extrusion process. The new crystalline structure revealed by microstructural investigation enhances the thermal properties of PLLA/PDLA film. It was found that the extrusion process had the influence on the thermal degradation temperature. Thermal treatment during the extrusion process probably leads to the polymer partial degradation. Therefore, a decrease of the degradation temperature is observed. The extrusion temperature has significant influence on the stereocomplexation, as the pure and stable sc PLA crystals were obtained at  $T_{ext} = 220$  °C and  $t_{ext} = 15$  min. In addition, the amorphous phase of sc PLA obtained by extrusion

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process relaxes independently from sc crystalline structure. However, the obtained samples were extremely mechanically fragile for further analysis.

The influence of the ratio of homopolymers was studied resulting in the pure sc PLA crystals formation in some cases. The optimum conditions of solution casting method were established as the pure sc PLA crystals were obtained only at ratio of 50:50 and 60:40 with the polymer concentration of 3 g/dL. It was found that solution casting has no influence on the thermal degradation temperature. However, the obtained pure sc PLA crystals were not stable as the appearance of the  $\alpha$  crystals was observed during 2<sup>nd</sup> heating scan.

In order to understand the properties of sc PLA, the further advance analysis will be carried out for the sc PLA sample obtained by solution casting (50:50 ratio of PLLA:PDLA, d= 3 g/dL). The amorphous phase species of sc PLA (obtained by solution casting and thermally molded-quenched according to the section 2.1.2.2.) will be detailed and discussed in the next chapter.

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**CHAPTER 4. MOLECULAR DYNAMIC STUDY IN AMORPHOUS STEREOCOMPLEX  
POLYLACTIDE**

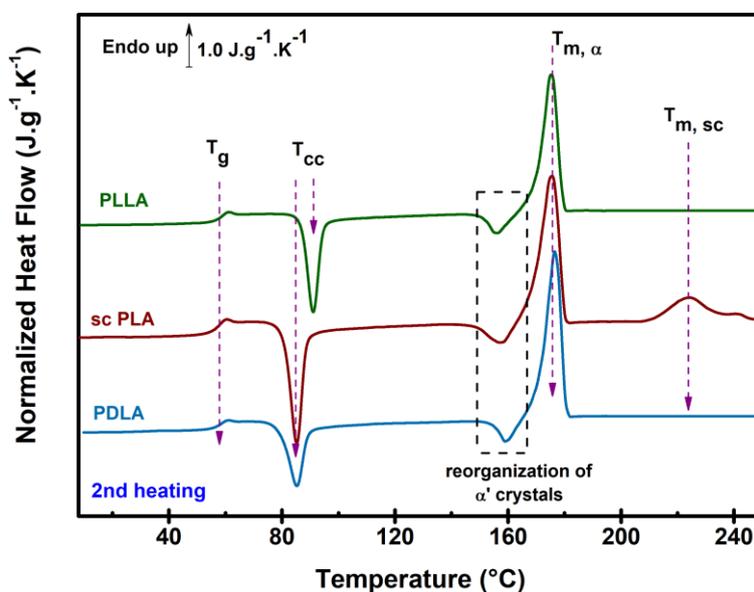
In this chapter, the influence of the tacticity on molecular dynamics of sc PLA's amorphous phase will be presented. In order to study the molecular dynamics the results of Fast Scanning Calorimetry (FSC) and Dielectric Relaxation Spectroscopy (DRS) were correlated. The combination of these two techniques allowed us to investigate the primary and secondary segmental relaxations of sc PLA and to compare them with the relaxations of amorphous homopolymers, PLLA and PDLA. In addition, different approaches (namely, fictive temperature, aging process, free volume model) were used in order to determine the amorphous phase properties.

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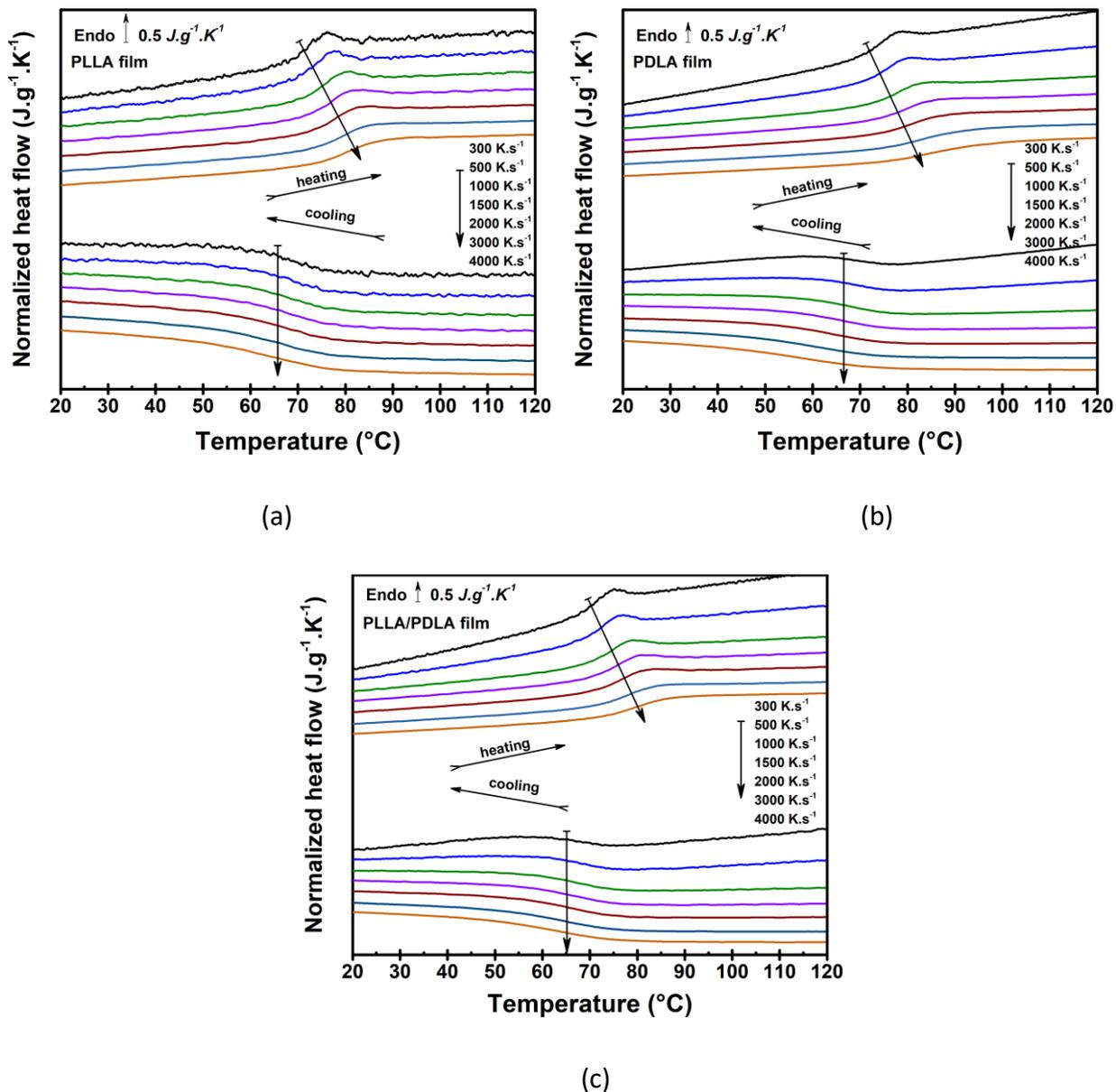
#### 4.1 Investigation of the amorphous phase properties through FSC

The glass transition temperature ( $T_g$ ) is the most important parameter for the structural relaxation analysis. Therefore, the evolution of  $T_g$  as a function of scanning rate was studied by means of the fictive temperature ( $T_f$ ) concept (presented in section 2.2.4.1) [1]. During the measurements special attention is paid to the sensitivity of the heat flows of FSC curves because of its dependence on the sample mass and scanning rate [2]. The obtained sample mass is equal to  $45 \pm 3$  ng according to the  $\Delta C_p = 0.46 \text{ Jg}^{-1}\text{K}^{-1}$  and  $0.54 \text{ Jg}^{-1}\text{K}^{-1}$  for homopolymers and sc PLA, respectively, measured by DSC (Fig 4.1), by comparing the value of the heat capacity step ( $\Delta C_p$ ) of wholly amorphous samples [3].



**Figure. 4. 1.** DSC normalized heat flow of the wholly amorphous PLLA, PDLA and sc PLA samples.

At the beginning samples were studied at temperatures from  $-50$   $^{\circ}\text{C}$  to  $270$   $^{\circ}\text{C}$  with scanning rate  $|\beta_c| = \beta_h$  ranging from  $300 \text{ Ks}^{-1}$  up to  $4000 \text{ Ks}^{-1}$  to determine the glass transition temperature. The normalized heat flows curves obtained by FSC are presented in Fig 4.2. The sensitivity of the signal becomes too low when decreasing the scanning rate from  $4000 \text{ K.s}^{-1}$  to  $300 \text{ K.s}^{-1}$  (Fig 4.2). Nevertheless, no enthalpy relaxation and thermal lag are observed during cooling. However, there are endothermic peaks which correspond to the enthalpy relaxation. Besides, shifts of the glass transition associated to the thermal lag can be observed during heating for each scanning rate.



**Figure. 4. 2.** FSC normalized heat flow of the wholly amorphous samples as a function of temperature upon heating and cooling at equivalent scanning rate  $|\beta_c| = \beta_h$  ranging from  $300 \text{ K.s}^{-1}$  up to  $4000 \text{ K.s}^{-1}$ : (a) PLLA, (b) PDLA, (c) sc PLA film samples. Arrows are given to follow the glass transition region.

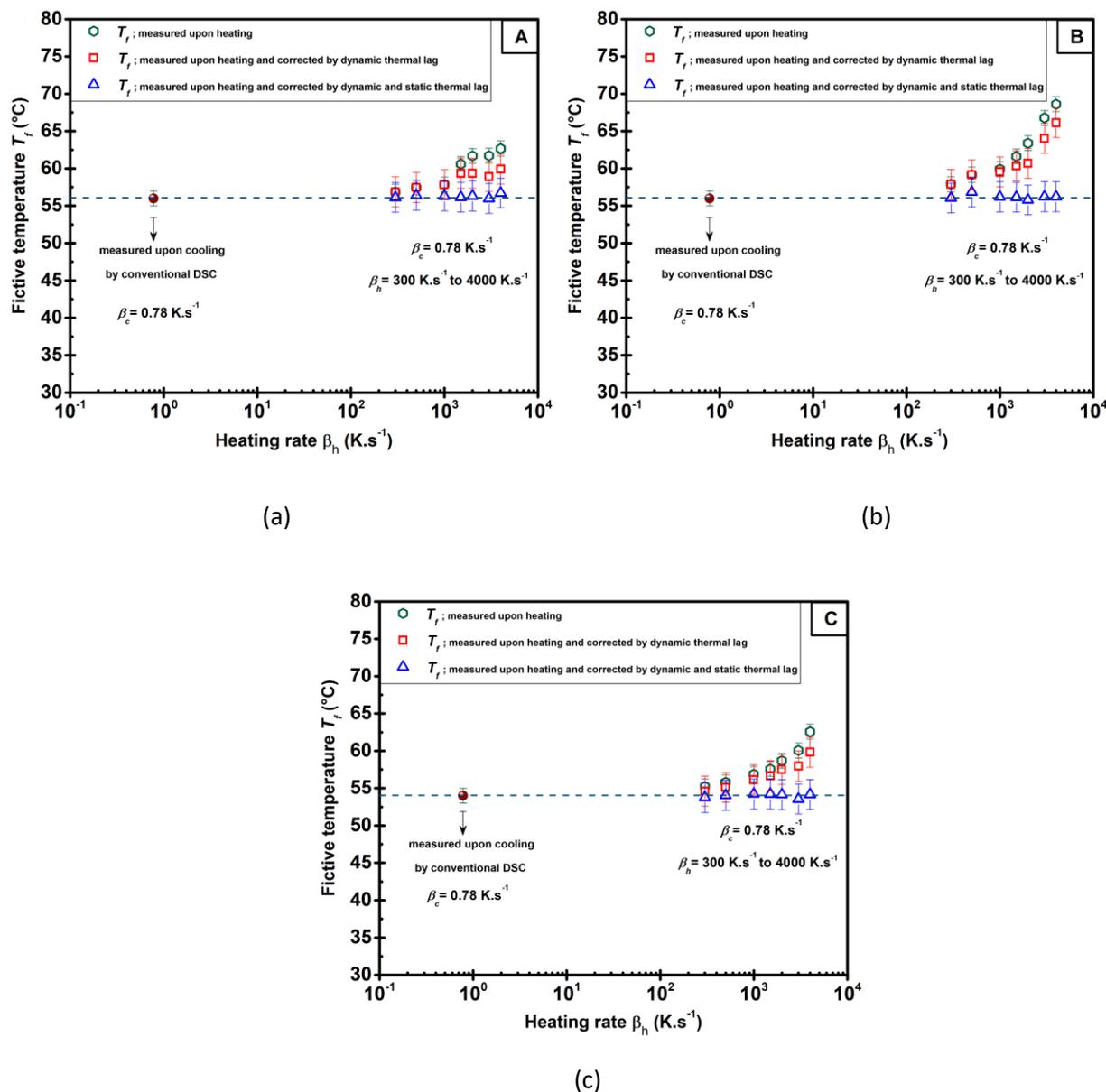
#### 4.1.1 The glass transition temperature and thermal lag corrections

The thermal lag obtained from FSC curves between heating and cooling is shown in Fig 4.2. Therefore, the thermal lag correction was carried out in order to improve the accuracy of the  $T_f$  investigation as detailed in section 2.2.4.2. The temperature corrections are done as proposed by Schawe [4] (values are presented in Appendix Chap 4 (A. 4.1)). The  $T_f$  and total

thermal lag were determined for wholly amorphous samples at scanning rate of seven decades – from  $300 \text{ K.s}^{-1}$  up to  $4000 \text{ K.s}^{-1}$ . At the rate of  $|\beta_c| = \beta_h = 300 \text{ K.s}^{-1}$ , no dynamic thermal lag  $\delta T_D$  was observed for PLLA and PDLA, while it was  $0.6 \text{ }^\circ\text{C}$  for sc PLA. The  $\delta T_D$  increases with the scanning rate increase, and it reaches nearly  $3 \text{ }^\circ\text{C}$  for  $|\beta_c| = \beta_h = 4000 \text{ K.s}^{-1}$  for all samples. On the contrary, a static thermal lag  $\delta T_S$  exists even at the lower scanning rates due to the sample thickness, and it increases with the increase of scanning rate. The calculated  $T_f$  upon heating increases with the scanning rate, while the calculated  $T_f$  upon cooling remains equal  $65 \pm 1 \text{ }^\circ\text{C}$  for PLLA and sc PLA, and  $66 \pm 1 \text{ }^\circ\text{C}$  for PDLA. The difference between the lowest and the highest values of the fictive temperature measured from heating is observed from  $5 \text{ }^\circ\text{C}$  up to  $10 \text{ }^\circ\text{C}$ . The average values of the  $T_f$  measured during cooling should match well with the average values of the corrected  $T_f$  from heating. Taking into account advantages of the  $\delta T_D$  and  $\delta T_S$  calibrations, the measured fictive temperatures from heating are corrected for each scanning rate (detailed in A. 4.1 and A. 4.2). The corrected  $T_f$  shows independent behavior from the scanning rate, and the average values are  $65 \pm 2$ ,  $66 \pm 2$ ,  $65 \pm 2 \text{ }^\circ\text{C}$  for PLLA, PDLA and sc PLA, respectively. In addition, Schawe purposed that the static effect can be prevented by the sample thickness of  $10 \text{ }\mu\text{m}$  or lower and, thus, the thermal lag occurs only due to the dynamic temperature gradient in case of the relatively lower scanning rates, i.e.  $< 100 \text{ K.s}^{-1}$  [4]. However, in this research high scanning rates were investigated in order to measure  $T_f$  during heating and cooling, thus  $\delta T_S$  might also be considered for samples with thickness lower than  $10 \text{ }\mu\text{m}$ . Therefore, the estimation of the  $T_f$  was carried out one more time for the same cooling rate and different heating rates. Since the  $T_f$  and/or  $T_g$  depend on the applied cooling rate, the same cooling rate ( $0.78 \text{ K.s}^{-1}$ ) is applied to all samples to vitrify, thus the unique  $T_f$  is measured during heating [5]. As the results of the vitrification, the calculated and corrected  $T_f$  by FSC must be equal the  $T_f$  measured upon cooling at  $0.78 \text{ K.s}^{-1}$  by DSC. The evolution of the measured  $T_f$  shows that when the heating rate increases, the heat transfer delay is observed in the samples and the  $T_f$  of the samples increases in the temperature range from  $54 \text{ }^\circ\text{C}$  up to  $69 \text{ }^\circ\text{C}$  (Fig 4.3). The  $T_f$  measured by FSC upon heating without correction,  $T_f$  corrected from  $\delta T_D$  and  $T_f$  corrected from  $\delta T_L$  are plotted in Fig 4.3. The  $T_f$  of wholly amorphous samples measured upon cooling by conventional DSC is also depicted by dark red sphere. Due to the straight line

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signal upon cooling on FSC, the  $T_f$  upon cooling was determined from the signal on conventional DSC as  $56 \pm 1$  °C for PLLA and PDLA,  $54 \pm 1$  °C for sc PLA as shown in Fig 4.3.



**Figure 4. 3.** Evolution of the  $T_f$  as a function of the heating rate  $\beta_h$  from  $300 \text{ K.s}^{-1}$  to  $4000 \text{ K.s}^{-1}$  investigated after vitrified the samples at  $|\beta_c| = 0.78 \text{ K s}^{-1}$ : (a) PLLA, (b) PDLA and (c) sc PLA. Dash line is average from blue empty up triangle (FSC) and dark red sphere (DSC).

Then, the values of  $T_f$  obtained by DSC were compared with the values of the corrected  $T_f$  from FSC. Only the  $\delta T_D$  was applied to the  $T_f$  to observe the necessity of the  $\delta T_S$  correction. As one can see, the  $T_f$  is still dependent on the heating rate after  $\delta T_D$  subtraction (Fig 4.3).

Therefore, the  $\delta T_D$  is not convenient to complete  $\delta T_L$  correction in case of high scanning rates. When  $\delta T_S$  was subtracted from the measured  $T_f$  as well as  $\delta T_D$ , the non-dependency of the  $T_f$  on heating rate and the matching between the corrected  $T_f$  from heating and the  $T_f$  measured upon cooling were observed (Fig 4.3).

Consequently, from the results of the thermal lag corrections, it may be noted that when the same cooling rate is applied, the same  $T_g$  and/or  $T_f$  is measured. It should be also mentioned that no significant difference of the  $T_f$  could be noted between the three polymers. On the other hand, the same values of the  $T_f$  were determined whatever the high cooling rates. Similar results were found for PLA-based polymers and its derivatives [3]. However, the difference between the  $T_f$  measured after cooling at  $|\beta_c| = 0.78 \text{ K}\cdot\text{s}^{-1}$  and  $|\beta_c| = 1500 \text{ K}\cdot\text{s}^{-1}$  is observed. Thus, this fact means that there is a critical cooling rate to obtain the unique glasses. Due to such difference between the values of  $T_f$  upon cooling at  $|\beta_c| = 0.78 \text{ K}\cdot\text{s}^{-1}$  and  $|\beta_c| = 1500 \text{ K}\cdot\text{s}^{-1}$ , the cooling rate dependency of  $T_f$  is well defined. Therefore, a lack of the applied cooling rate dependence of the fictive temperature can be observed in glass-forming systems above certain and/or critical scanning rate during FSC measurements [6, 7].

#### **4.1.2 The fragility index $m$**

The cooling rate dependence of the  $T_f$  was studied and it was fitted by the VFTH equation (Eq 2.15) (Appendix Chap 4 (A. 4.3)) to determine the dynamic fragility index  $m$  by VFT equation (Eq 2.16 and Eq 2.17) [5, 8-10]. In the  $m$  calculation,  $T_g$  is taken as the  $T_f$  determined at a cooling rate of  $0.78 \text{ K}\cdot\text{s}^{-1}$  as 329 K and 327 K for homopolymers and sc PLA, respectively. According to the obtained results (Table 4.1), the calculated values of the dynamic fragility  $m$  for studied films are rather close taking into account the measurement precision. These values are in the range of the fragility values defined by Angell:  $16 \leq m \leq 200$ . The higher the value of  $m$ , the more fragile the system. So, all the samples can be defined as "fragile".

Sample	$B$ (K)	$T_V$ (K)	$M$
PLLA	$36 \pm 11$	$48 \pm 3$	$185 \pm 17$
PDLA	$37 \pm 8$	$48 \pm 3$	$190 \pm 11$
Sc PLA	$73 \pm 22$	$41 \pm 4$	$142 \pm 44$

**Table 4. 1.** The VFTH fitting parameters and the dynamic fragility index of studied samples.

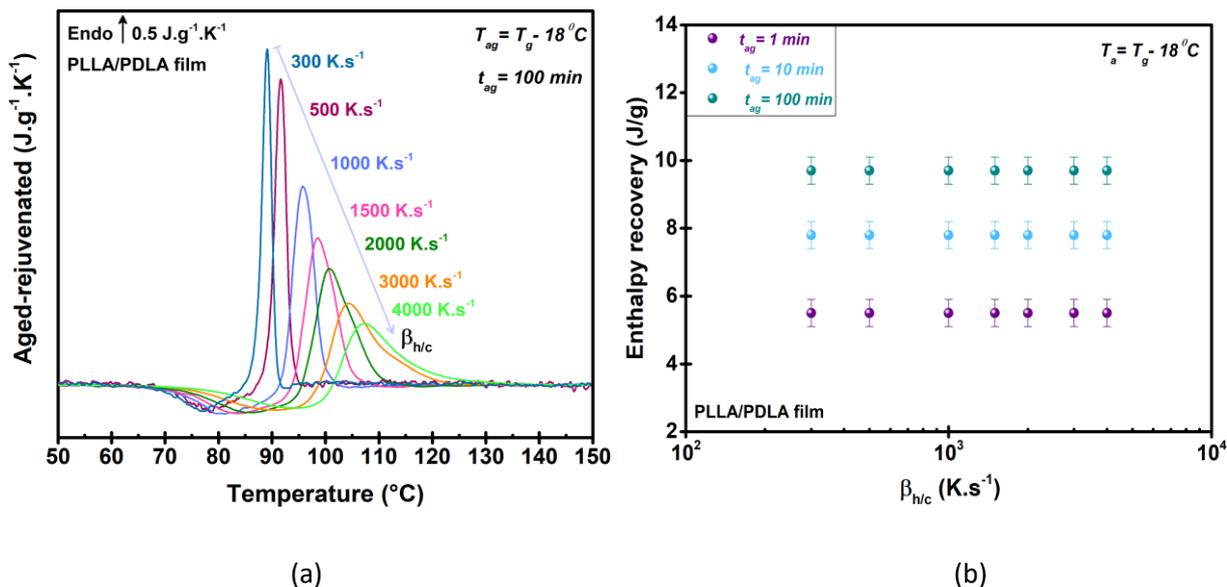
## 4.2 Structural relaxation dynamics through physical aging

The measured and corrected  $T_f$  values were determined to estimate the aging temperature ( $T_{ag}$ ). The determined constant values of  $T_f$  at all scanning rates exhibit unique glass formation. Due to the estimation of unique glass obtained by higher cooling rate, the scanning rate dependence of the physical aging was investigated.

### 4.2.1 Influence of the scanning rate on physical aging

Physical aging was carried out at the  $T_{ag}$  calculated according to the corrected  $T_g$  which are 66 °C for PDLA and 65 °C for PLLA and sc PLA after a cooling at 1500 K.s<sup>-1</sup>, as shown in section 2.2.4.3. The recovery enthalpy values were calculated and compared for each scanning rate and aging time ( $t_{ag}$ ) for all the samples (detailed in section 2.2.4.3) [11]. The enthalpy recovery for each samples (PLLA, PDLA and sc PLA) at the scanning rate range for  $t_{ag} = 1$  min, 10 min and 100 min has similar behavior. Therefore, only the results of aging at the scanning rate  $|\beta_c| = \beta_h$  ranging from 300 K.s<sup>-1</sup> up to 4000 K.s<sup>-1</sup> are shown as an example (Fig 4.4).

As one can see from Fig 4.4a, the peaks are shifted towards higher temperatures with the increase of the scanning rate due to the smearing effects. In addition, as shown in Fig 4.4b, the values of enthalpy recovery are independent on the scanning rate, but dependent on the  $t_{ag}$ , as average values of  $5.5 \pm 0.4$  J/g,  $7.8 \pm 0.3$  J/g, and  $9.7 \pm 0.4$  J/g for 1 min, 10 min, and 100 min, respectively, are obtained. Thus, not only the fictive temperature shows independence on the scanning rate, but also the enthalpy recovery is independent from scanning rate above the critical cooling rate. Therefore, for further physical aging analysis to compare the structural relaxation of amorphous PLLA, PDLA and sc PLA, the scanning rate of  $|\beta_c| = \beta_h = 1500$  K.s<sup>-1</sup> was chosen due to the less signal-noise ratio.

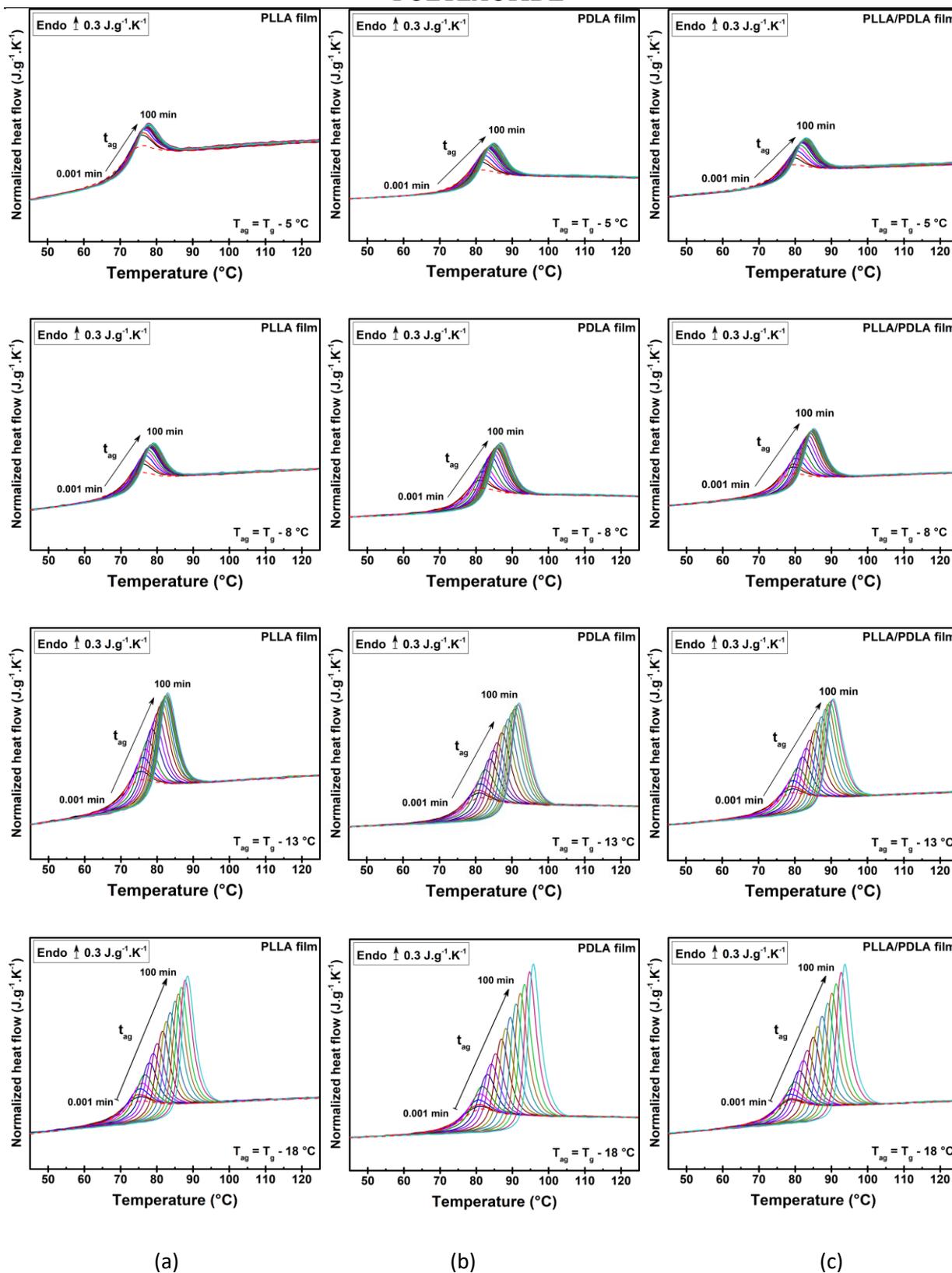


**Figure 4. 4.** Aging of sc PLA at  $T_{ag} = T_g - 18^\circ\text{C}$  at the scanning rate range  $|\beta_c| = \beta_h$  from  $300\text{ K}\cdot\text{s}^{-1}$  up to  $4000\text{ K}\cdot\text{s}^{-1}$ . (a) Normalized heat flow subtraction for  $t_{ag} = 100\text{ min}$ . (b) Values of the enthalpy recovery as a function of  $\beta_{h/c}$  for  $t_{ag} = 1\text{ min}$ ;  $10\text{ min}$ ;  $100\text{ min}$ .

#### 4.2.2 Physical aging at four different aging temperature

The physical aging was carried out at the  $T_{ag}$  chosen as  $T_{ag} = T_f - 5^\circ\text{C}$ ,  $T_{ag} = T_f - 8^\circ\text{C}$ ,  $T_{ag} = T_f - 13^\circ\text{C}$  and  $T_{ag} = T_f - 18^\circ\text{C}$  with the  $t_{ag}$  ranging from  $0.001\text{ min}$  to  $100\text{ min}$  for all the samples. The enthalpy recovery of the structural relaxation of amorphous samples was investigated to define the chirality impacts on the relaxation. Physical aging of polymers was widely studied and it was reported that quenching or higher cooling rates allowed us to accelerate the aging and the enthalpy recovery of materials to reach equilibrium on laboratory scale [3, 12-15]. Fig 4.5 presents the normalized heat flows of all the aged samples from  $0.001\text{ min}$  up to  $100\text{ min}$  obtained by FSC at  $|\beta_c| = \beta_h = 1500\text{ K}\cdot\text{s}^{-1}$  for the  $T_{ag}$  below the respective glass transition temperature. The red dash line corresponds to the rejuvenated curve. For all the curves, endothermic relaxation peaks are superimposed to the glass transition and shift towards higher temperature with the increasing  $t_{ag}$ , as usually observed for physical aging [3, 12-15]. The endothermic peaks are intense and broadened, thus, the lower the  $T_{ag}$ , the higher the enthalpy recovery. Fig 4.6 shows the decline of the enthalpy during the physical aging for all studied  $T_{ag}$ .

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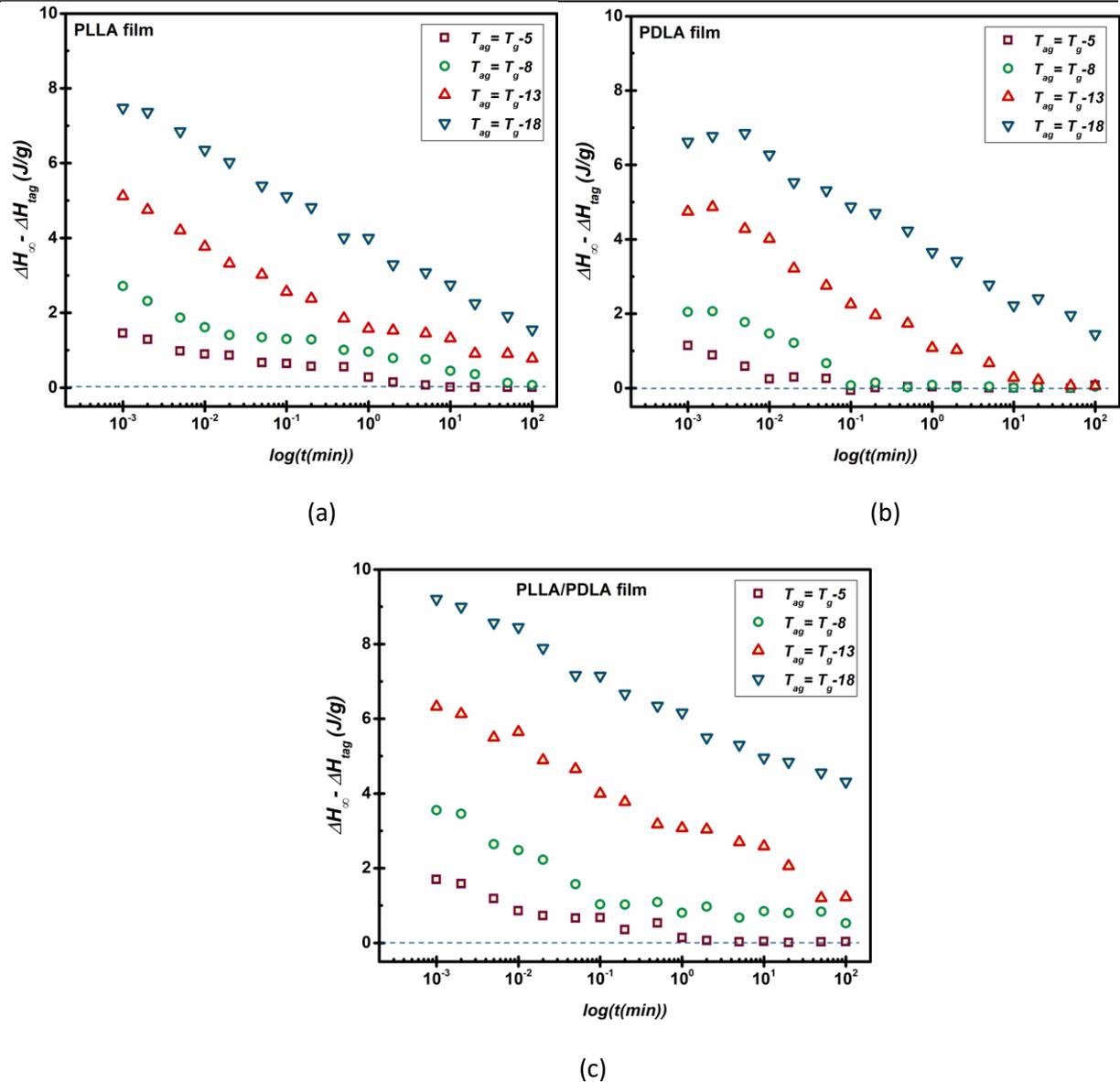
**Figure 4. 5.** Aged amorphous (a) PLLA, (b) PDLA and (c) sc PLA films examined by FSC. Graphs on the same line represent the same  $T_{ag}$ .

The difference between the total enthalpy loss  $\Delta H_{\infty}$  and the enthalpy recovery at a given aging time  $\Delta H_{tag}$  shows different behaviors according to the  $T_{ag}$ . Despite the different structure of samples, similarities are observed for the same  $T_{ag}$  and  $t_{ag}$ . The kinetic decay of the enthalpy recovery during the physical aging is similar for different samples. A rapid kinetics of physical aging was observed as the equilibrium was reached after 1 min at 5 °C below  $T_g$ . Besides the rapid kinetics, FSC also allows us to analyze few ng sample as a reference of the bulk sample in shorter aging time such as 0.001 min due to the formation of glass with high fictive temperature with very short relaxation time [16]. The enthalpy recovery is guided by a single step decay and the classical stretched tendency is observed similar to other glassy polymers [16-18].

As depicted in Fig 4.6, the enthalpy recovery exhibits a single step decay towards equilibrium and the glass completely releases its excess of energy in the range of aging time at the highest aging temperature. Such result is opposite to the assumption proposed by Gomez Ribelles et al. [19, 20] based on DSC measurements. The authors assumed that thermodynamic equilibrium cannot be reached because of the steric hindrance and molecular mobility induced during structural relaxation. In addition, Hutchinson et al. [21] reported that a certain fraction of the glass-forming liquid was vitrified towards the glass transition. Therefore, the slow-process (i.e. the vitrified fraction) and the fast-process (i.e. the non-vitrified fraction inhibit) to reach the thermodynamic equilibrium. FSC allows obtaining glasses with high enthalpy and reaching the thermodynamic equilibrium. The lowest aging temperatures exhibits also single step decay, but does not allow reaching the equilibrium at the investigated aging time.

Although all the samples have the similar behavior (i.e. a single step decay), in order to compare the relaxation kinetics of all the amorphous samples, the enthalpy recovery and the master curves of normalized enthalpies are investigated and presented in Fig 4.7.

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**Figure 4. 6.** Time evolution of the difference between the  $\Delta H_\infty$  and the  $\Delta H_{tag}$  for the aged amorphous (a) PLLA, (b) PDLA and (c) sc PLA as a function of  $T_{ag}$  with the aging time ranging from 0.001 min to 100 min at  $|\beta_c| = \beta_h = 1500 \text{ K}\cdot\text{s}^{-1}$ .

The difference in the calculated total enthalpy loss  $\Delta H_\infty$  is caused by the amplitude of the heat capacity step of the samples ( $0.46$  and  $0.54 \text{ J}\cdot\text{g}^{-1}\cdot\text{K}^{-1}$  for homopolymers and sc PLA, respectively). As shown in Fig 4.7a, the physical properties are dependent on the logarithm of the  $t_{ag}$ . Therefore, the aging rate can be determined as the slope of the linear fit and corresponds to the enthalpy change towards equilibrium. One can noticed from the Fig 4.7a that all the samples have the similar rate of enthalpy recovery ( $1.36 \pm 0.06 \text{ J/decade}$ ). Fig 4.7b presents superimposition of the master curve of normalized enthalpy recovery as a function

of the logarithm of the  $t_{ag}$ . The enthalpy recoveries are normalized by the infinite enthalpy loss as follows:

$$\Delta H_{normalized} = \frac{\Delta H_{\infty} - \Delta H_{recovery}}{\Delta H_{\infty}} \quad (4.1)$$

where  $\Delta H_{normalized}$  is the normalized enthalpy recovery,  $\Delta H_{recovery}$  is the measured enthalpy recovery of the aged samples, and  $\Delta H_{\infty}$  is the total enthalpy loss.

In addition, the Tool-Narayanaswamy-Moynihan (TNM) model is used to explain the evolution of the glass transition temperature related to the relaxation enthalpy and aging temperature ( $T_{ag}$ ) for an aging time ( $t_{ag}$ ) [11, 22]. The TNM model is used to fit master curves by assuming a specific expression for the temperature and structure dependence of the relaxation time  $\tau(T, T_g)$  [23] as follows:

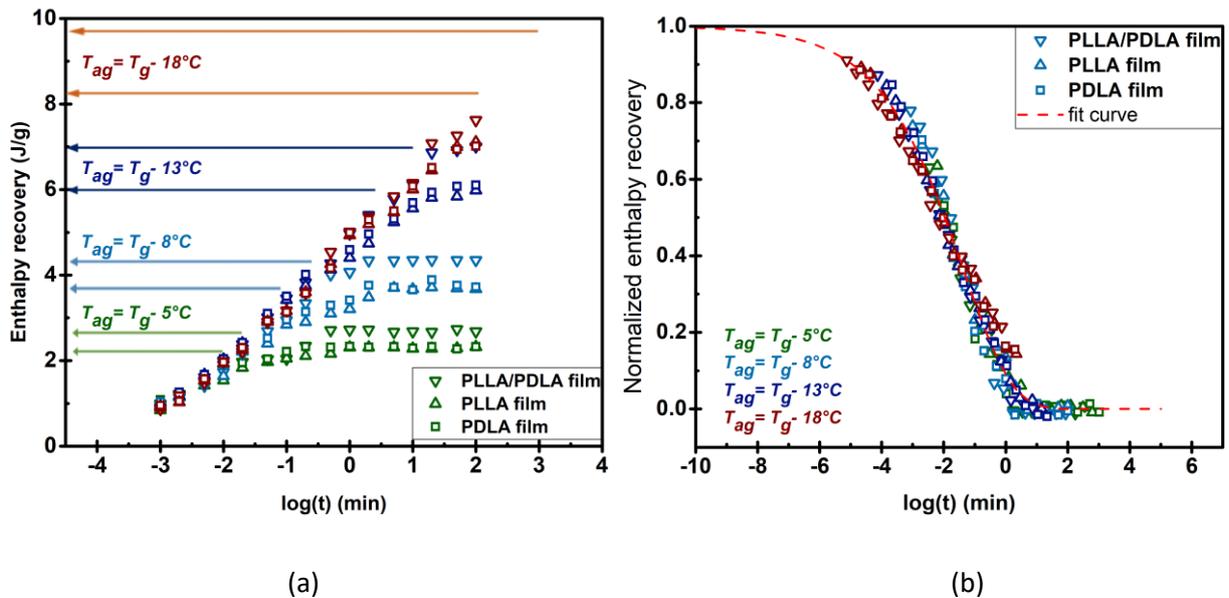
$$\tau(T_f, T) = A \exp \left\{ \frac{x \Delta H}{RT} + \frac{(1-x)\Delta H}{RT_f} \right\} \quad (4.2)$$

where  $T$  is a reference temperature well above the glass transition temperature,  $x$  is the nonexponential parameter determining the broadness of the relaxation spectrum. When these parameters are obtained by DSC analysis, so the shape of parameters corresponds to the behavior of the system in the out-of-equilibrium state. However, some inconsistencies were reported concerning the equilibrium behavior if the relaxation expressed for  $\tau(T, T_f)$  was not fitted correctly. On the contrary to the complexity of TNM model, in our case, the master curves of enthalpy recovery of all the samples can be fitted using a simplified approach, i.e. the Kohlrausch-Williams-Watts (KWW) function [24], as follows:

$$\phi(t) = e^{-(t/\tau_{KWW})^{\beta_{KWW}}} \quad (4.3)$$

where  $\Phi(t)$  is the correlation function,  $\beta_{KWW}$  ( $0 < \beta_{KWW} < 1$ ) is a stretch exponent, and  $\tau_{KWW}$  is the relaxation time for  $\beta_{KWW} = 1$ . As shown in Fig 4.7, the master curves of  $T_{ag} = T_g - 8$  °C were taken as a reference temperature to impose all the master curves and the other curves were shifted over the reference curve with the shift factor  $x = 0.4$  for  $T_{ag} = T_g - 5$  °C and  $x = -0.2$  for  $T_{ag} = T_g - 13$  °C and  $T_{ag} = T_g - 18$  °C. In this case, KWW function is sufficient to fit all the curves

within the single relaxation time  $\tau$  which is determined as 0.04 s according to the reference temperature. The kinetics of decays of the enthalpy recovery of all the samples during the physical aging is exactly the same. No influence of the tacticity on the structural relaxation is observed as a result of kinetic investigation.



**Figure 4. 7.** (a) Time evolution of the enthalpy recovery  $\Delta H_t$  for aged amorphous PLLA (up triangle), PDLA (square) and sc PLA (down triangle) films at different  $T_{ag}$ . The arrows show the values of  $\Delta H_\infty$  for homopolymers and sc PLA. (b) Master curves of the enthalpy recovery normalized by the infinite enthalpy loss. Red dash line corresponds to the KWW fitting (Eq 4.3).

### 4.3 Segmental relaxation investigations by DRS

The molecular dynamics of all the amorphous samples was investigated by means of dielectric relaxation spectroscopy (DRS). The  $\alpha$ -relaxation parameters like the dielectric strength, the relaxation time and the fragility index of all samples were examined to combine the results with the results of FSC. The cooperativity of molecular mobility is presented in the notion of the dynamically correlated number  $N_c$  and the cooperativity degree  $N_\alpha$ .

#### 4.3.1 The $\alpha$ -relaxation process

The experimental raw data obtained from dielectric relaxation spectroscopy measurements are presented in 3-D illustration in Fig 4.8. The dielectric loss  $\epsilon''$  is measured as a function of temperature and frequency. The dielectric spectra of all samples display

similar characteristics in the same temperature range towards the glass transition. The  $\alpha$ -relaxation peak which is the dielectric response of the glass transition, broadens and shifts towards higher temperature as the frequency increases. The conductivity phenomenon which is related to the charge transport is also observed at high temperature and low frequency. The dielectric relaxation spectra can be characterized by a “step-like” decrease of  $\epsilon'$  and a peak of  $\epsilon''$  in the frequency range at an isothermal temperature. The main parameters to characterize dielectric properties can be determined from both  $\epsilon'$  and  $\epsilon''$  versus frequency plot. As an example, the relaxation time  $\tau_{max}$  can be determined from the maximum value of frequency  $f_{max}$  where  $\epsilon''$  reaches its maximum value at a certain temperature  $T$ . The measured complex permittivity signals ( $\epsilon^*$ ) associated with the primary ( $\alpha$ -relaxation) and secondary ( $\beta$ -relaxation) relaxation were fitted by the empirical Havriliak-Negami (HN) function (Eq 4.4) to complete quantitative analysis of the dielectric spectra in the frequency domain [25]:

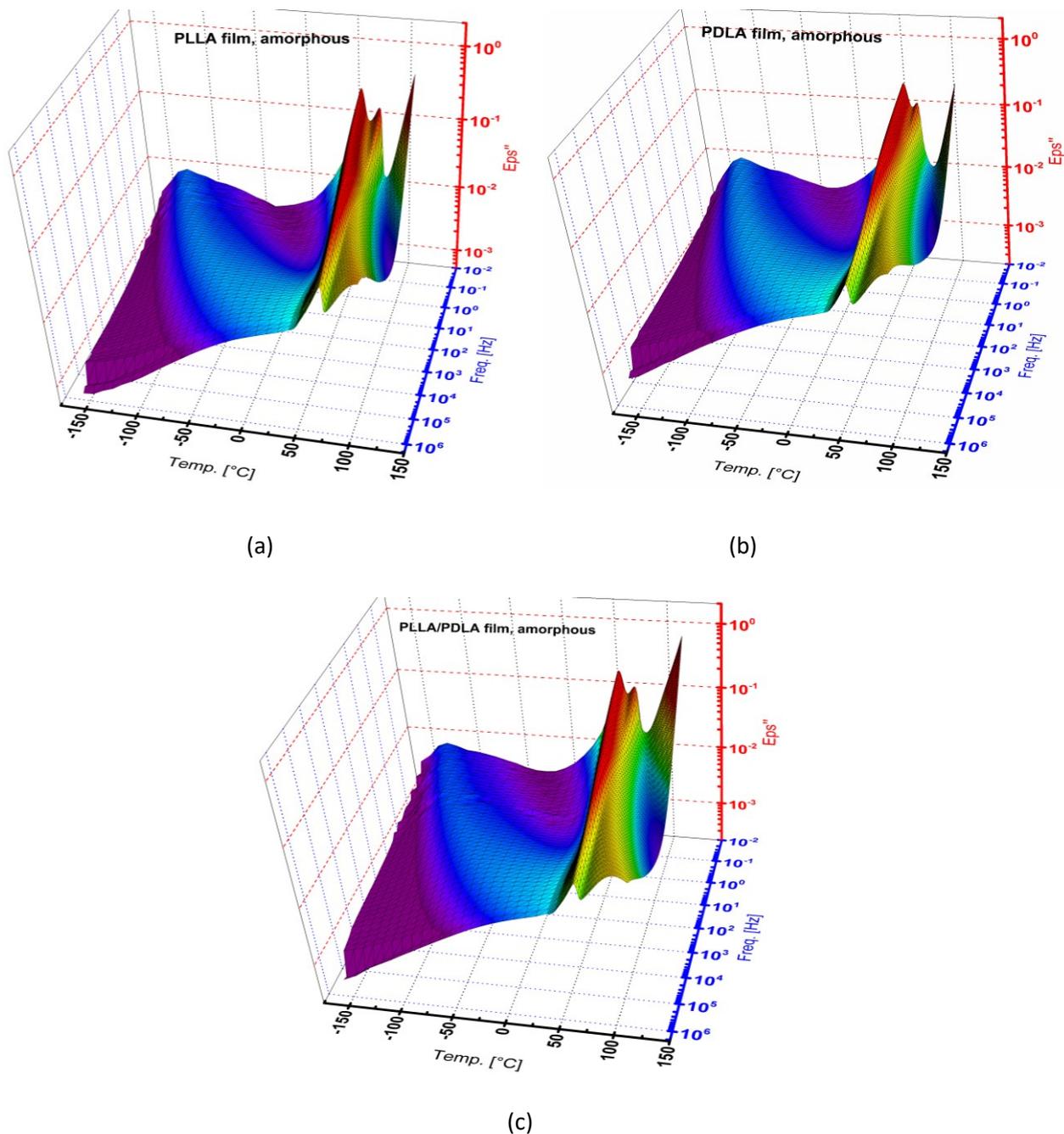
$$\epsilon^*(\omega) = \epsilon_{\infty} + \frac{\Delta\epsilon}{[1 + (i\omega\tau_{HN})^{\alpha_{HN}}]^{\beta_{HN}}} \quad (4.4)$$

The HN function allows to fit the real ( $\epsilon'(\omega)$ ) and the imaginary ( $\epsilon''(\omega)$ ) components of the complex dielectric permittivity ( $\epsilon^*(\omega)$ ) by the following equations:

$$\epsilon'(\omega) = \epsilon_{\infty} + \Delta H_{HN} \frac{\cos(\beta_{HN}\varphi_{HN})}{\left[1 + 2 \sin\left(\frac{\pi(1-\alpha_{HN})}{2}\right)(\omega\tau_{HN})^{\alpha_{HN}} + (\omega\tau_{HN})^{2\alpha_{HN}}\right]^{\beta_{HN}/2}} \quad (4.5)$$

$$\epsilon''(\omega) = \Delta H_{HN} \frac{\sin(\beta_{HN}\varphi_{HN})}{\left[1 + 2 \sin\left(\frac{\pi(1-\alpha_{HN})}{2}\right)(\omega\tau_{HN})^{\alpha_{HN}} + (\omega\tau_{HN})^{2\alpha_{HN}}\right]^{\beta_{HN}/2}} \quad (4.6)$$

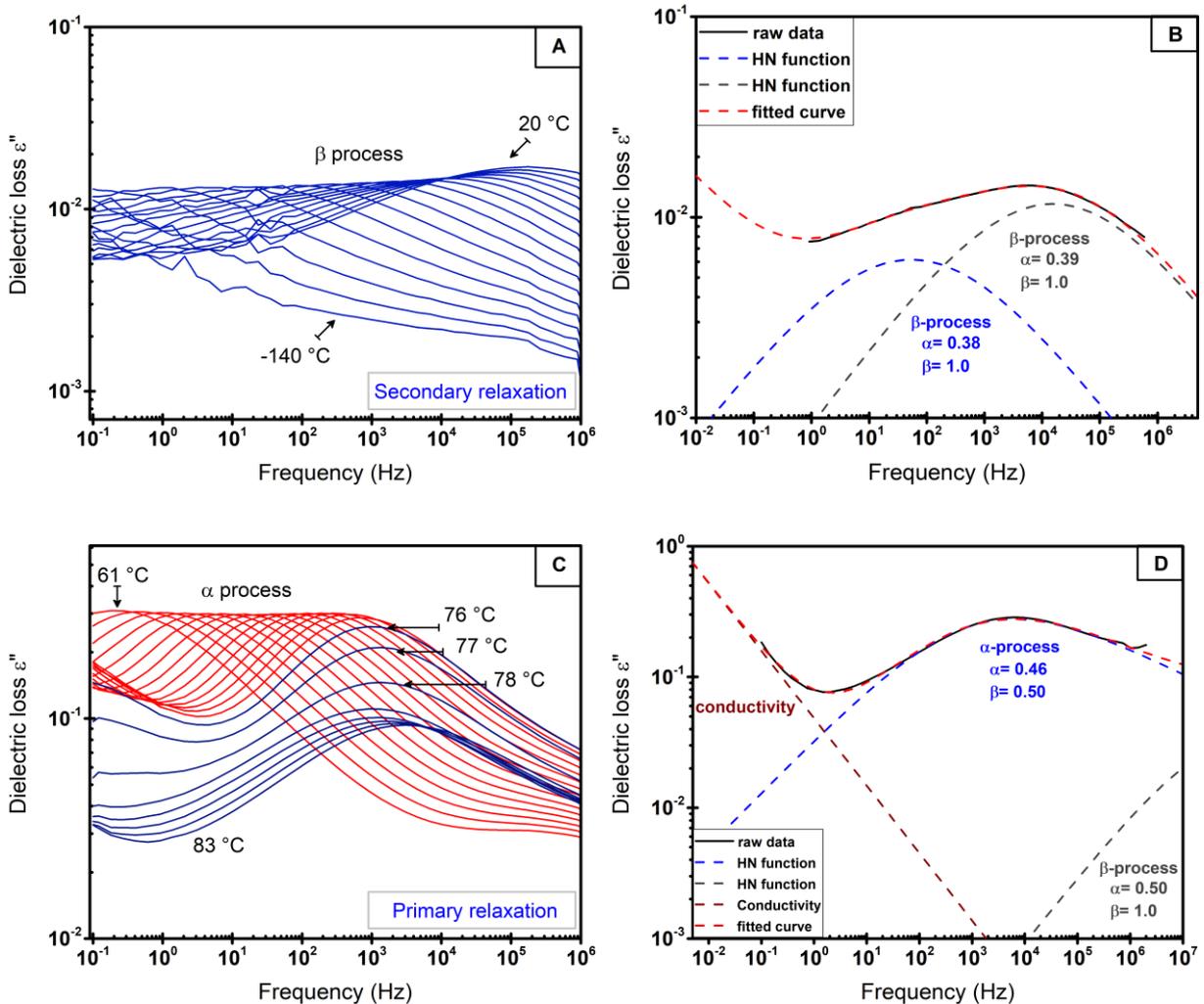
where  $\omega$  is the angular position ( $\omega = 2\pi f$ ),  $\epsilon_{\infty}$  is the unrelaxed dielectric permittivity,  $\Delta\epsilon_{HN}$  is the relaxation strength,  $\tau_{HN}$  is a characteristic relaxation time and  $\alpha_{HN}$  and  $\beta_{HN}$  are shape parameters describing the symmetric and asymmetric broadening factor of the dielectric spectra, respectively. In order to improve the consistency of the fit results, the fitting has been carried out for both real and imaginary parts of the complex signals.



**Figure 4. 8.** Dielectric loss ( $\epsilon''$ ) versus frequency and temperature for all studied amorphous samples: (a) PLLA, (b) PDLA and (c) sc PLA.

The isothermal dielectric spectra of amorphous sc PLA are shown as a function of frequency for different temperatures in Fig 4.9A and Fig 4.9C as an example of DRS analysis. All the amorphous samples show similar variations during relaxation investigations. It should be noted that all samples show two complex relaxation processes. The first one is the

secondary relaxation ( $\beta$ -process) observed at low frequencies and temperatures (Fig 4.9A). This relaxation shifts towards higher frequency range as the temperature increases. The second one is the primary relaxation ( $\alpha$ -process) which can be clearly seen in higher temperature range (Fig 4.9C).



**Figure 4. 9.** Imaginary parts of the complex dielectric permittivity ( $\epsilon''$ ) versus frequency of amorphous sc PLA film: (A) in the glassy state at temperature  $T < T_g$ , (B) clarification of dielectric spectra fitting for secondary relaxation ( $\beta$ -process), (C) in the liquid state at temperature  $T > T_g$ , and (D) clarification of dielectric spectra fitting for primary relaxation ( $\alpha$ -process) by HN complex functions.

The two HN complex functions are used to fit isothermal dielectric loss spectra with a conductivity contribution to investigate the relaxation phenomena. The HN shape parameters of both relaxations in amorphous sc PLA were shown in Fig 4.9B and 4.9D. All amorphous

samples have the same temperature dependence for the main relaxation processes, in which both  $\alpha$  and  $\beta$  relaxations shift to higher frequencies but keep the same amplitude as the temperature increases until the cold crystallization temperatures of amorphous samples. As shown in Fig 4.9C, there is a strong reduction in the amplitude of dielectric loss above  $T = 75^\circ\text{C}$  due to the cold crystallization during the measurements. Therefore, the analytical procedure of fitting was carried out until the first crystallization appeared. The demonstration of complex secondary relaxation with two contributions is supported by the results obtained for the other polyesters, such as poly(ethylene terephthalate) (PET) [26, 27], PLA [28] and poly(hydroxyalkanoates) (PHAs) [29]. In addition, Soccio et al. [30] reported that such a broad  $\beta$  relaxation phenomenon is caused by interactions between ester functional groups and the main polymer chain as well as stereocomplexation between homopolymers (PLLA and PDLA). In the case of poly(butylene 2,5-furanoate), the  $\beta$  relaxation has two processes which are a faster  $\beta_1$  relaxation due to the association of the mobile subunit with the C – O of the ester group and a slower  $\beta_2$  relaxation correlated to the connection between the aromatic ring and C = O of the ester group. In the case of stereocomplexation, such a classical broad  $\beta$  relaxation of polyesters maybe caused by similar interactions between the mobile subunit and the C – O of the ester group and/or the main chain and C = O of the ester group due to the stereoselective van der Waals forces. Therefore, the isothermal spectra of  $\beta$  relaxation were analyzed by two symmetrical HN functions, called Cole-Cole functions at different temperatures.

#### 4.3.1.1 The master curves and shape parameters: $\alpha$ and $\beta$

The shape parameters determined by the HN fits of the  $\alpha$ -relaxation process are used to define the symmetric and asymmetric broadening of the complex dielectric function [31]. The values of the shape parameters determined for all the samples are found to be less than 1 in accordance with literature [31]. By using the estimated values of  $\tau_{HN}$ ,  $\alpha_{HN}$ ,  $\beta_{HN}$  parameters, the relaxation time,  $\tau_{max} = (2\pi f_{max})^{-1}$  was calculated as [31]:

$$\tau_{max} = \tau_{HN} \times \left[ \frac{\sin\left(\frac{\alpha_{HN}\beta_{HN}\pi}{2+2\beta_{HN}}\right)}{\sin\left(\frac{\alpha_{HN}\pi}{2+2\beta_{HN}}\right)} \right]^{1/\alpha_{HN}} \quad (4.7)$$

The correlation function can be calculated using the HN relationship [32]:

$$f(\tau) = \frac{1}{\pi} \frac{\left(\frac{\tau}{\tau_{HN}}\right)^{\beta_{HN}\alpha_{HN}} \sin(\beta_{HN}\varphi)}{\left[1 + \left(\frac{\tau}{\tau_{HN}}\right)^{2\alpha_{HN}} + 2\left(\frac{\tau}{\tau_{HN}}\right)^{\alpha_{HN}} \cos(\pi\alpha_{HN})\right]^{\beta_{HN}/2}} \quad (4.8)$$

with

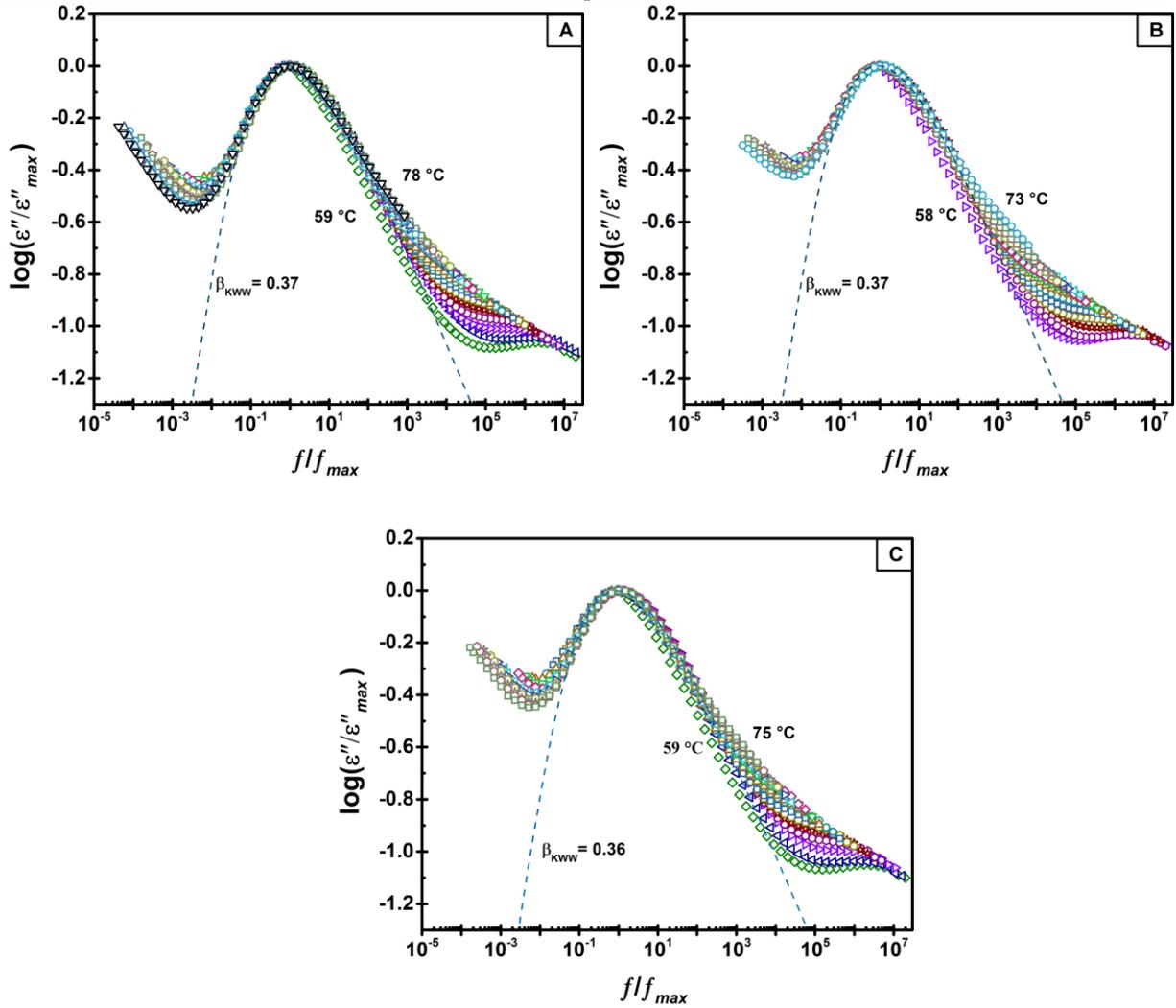
$$\varphi = \frac{\pi}{2} - \tan^{-1} \left[ \frac{\left(\frac{\tau_{HN}}{\tau}\right)^{\alpha_{HN}} + \cos(\pi\alpha_{HN})}{\sin(\pi\alpha_{HN})} \right] \quad (4.9)$$

where  $f(\tau)$  is the distribution function,  $\tau_{HN}$  is the HN relaxation time,  $\alpha_{HN}$  and  $\beta_{HN}$  are the HN shape parameters. Therefore, in term of the distribution of relaxation time, the correlation function  $\Phi(t)$  is expressed by [32]:

$$\Phi(t) = \int_0^{\infty} f(\tau) \exp^{-t/\tau} dt \quad (4.10)$$

In a certain time domain, the correlation function can be well described by the KWW stretch exponential function as presented in Eq 4.3 [33]. To express the time-scale required for molecular units to relax or move, the master plots of the dielectric loss for each isothermal spectrum are reported in Fig 4.10. The results show very good overlap of all the curves forming a single master curve. On the other hand, systematic deviation from the master curve is observed in the high frequency range for the  $\alpha$  relaxation. This kind of deviation can be explained by the contribution of noncooperative localized molecular mobility, such as  $\beta$  relaxation process, combined with the molecular mobility of localized molecules. Furthermore, such an interdependence of  $\alpha$  and  $\beta$  relaxations was reported also by Johari and Goldstein [34] and was investigated in glass-forming systems [35, 36]. While the temperature of measurement is increased, the impact of the  $\beta$  relaxation increases on the master curves. The non-Debye relaxation in the time range ( $t$ ) is empirically explained by the KWW function [24]. Each master curve was fitted by the KWW relationship (Eq 4.3) in order to calculate KWW stretching parameter  $\beta_{KWW}$ . The stretching parameter  $\beta_{KWW}$  allows comparing any asymmetrical broadening behavior of the relaxation process at high frequencies with the exponential decay which corresponds to a Debye relaxation with  $\beta_{KWW} = 1$ .

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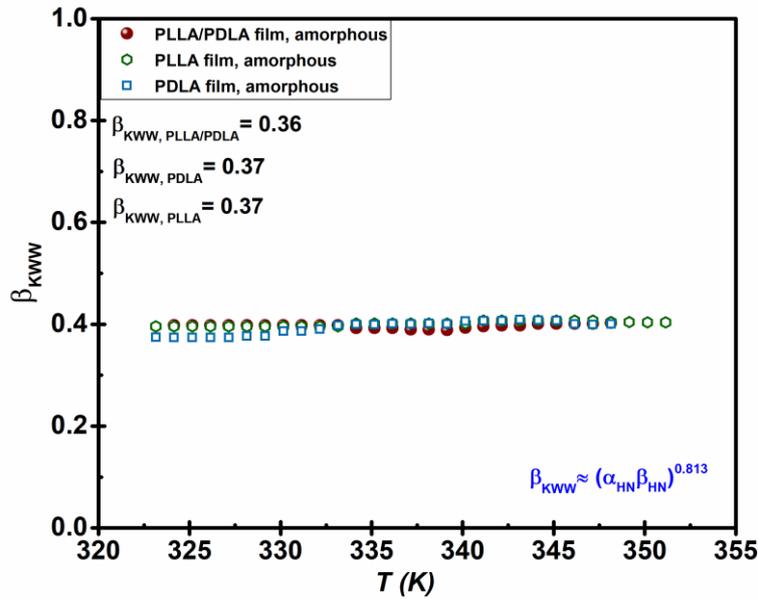
**Figure 4. 10.** Log-log plot of the master curves of amorphous (A) PLLA, (B) PDLA and (C) sc PLA samples. Light blue dash lines correspond to the KWW fits.

The HN parameters are correlated with the stretching parameter  $\beta_{KWW}$  as follows [31]

$$\beta_{KWW} = (\alpha_{HN}\beta_{HN})^{0.813} \quad (4.11)$$

The master curves were obtained by normalization of the spectra with the maximum of  $\epsilon''$  and its counterpart frequency at the different temperatures (from 58 °C to 78 °C with a step of 1 °C) and fitted by KWW function to graphically visualize the stretching parameter  $\beta_{KWW}$ . These master plots lead to show whether temperature has influenced on the shape of the structural relaxation process, which would represent different distributions of the relaxation times at different temperatures. As shown in Fig 4.11, the values of  $\beta_{KWW}$  are experimentally found to be  $0.37 \pm 0.02$  for PLLA and PDLA, and  $0.36 \pm 0.02$  for sc PLA. These values have a

good compliance with the values calculated by Eq 4.16:  $0.40 \pm 0.01$  for PLLA,  $0.39 \pm 0.01$  for PDLA and  $0.39 \pm 0.01$  for sc PLA. In addition, This result is in good agreement with the value reported in literature for PLA from DSC measurements,  $\beta_{KWW} \approx 0.35$  [37].



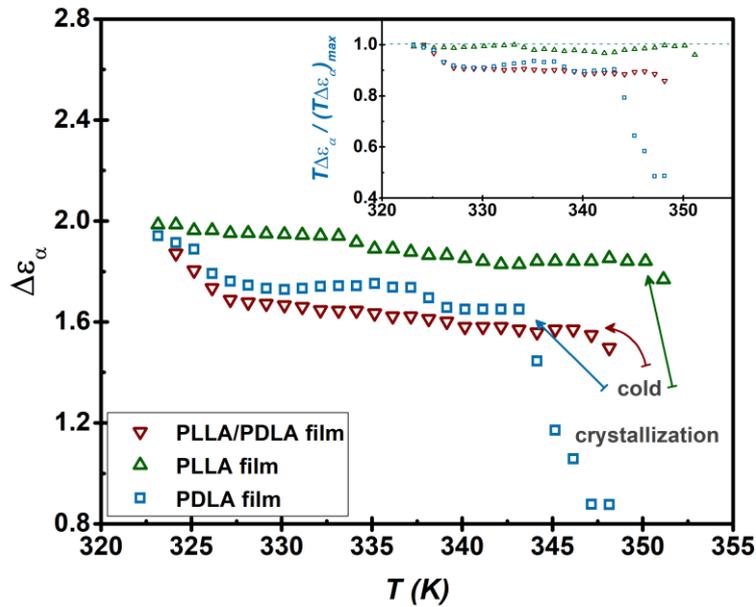
**Figure 4. 11.** Variation of the  $\beta_{KWW}$  parameter as a function of temperature for studied samples.

#### 4.3.1.2 Dielectric relaxation strength $\Delta\varepsilon_\alpha$

The dielectric relaxation strength  $\Delta\varepsilon_\alpha$  of  $\alpha$  relaxation recorded for all the isothermal measurements is obtained by fitting the experimental data with the HN function. In addition, as reported by Onsager, Fröhlich, and Kirkwood [31], the values of  $\Delta\varepsilon_\alpha$  depend on several parameters and are also calculated by the generalized form of the Debye's theory as follows:

$$\Delta\varepsilon_\alpha = \frac{1}{3\varepsilon_0} g_K \frac{\mu^2}{k_B T} \frac{N}{V} \quad (4.12)$$

where  $\varepsilon_0$  is the dielectric permittivity of vacuum,  $g_K$  is the Kirkwood correlation factor which corresponds to the short-range intermolecular interactions leading to specific static dipole-dipole orientations,  $\mu^2$  is the time-correlation function of the total dipole moment,  $k_B$  is the Boltzmann's constant,  $T$  is the temperature, and  $N/V$  is the volume density of dipoles involved in the relaxation process. Fig 4.12 presents the temperature dependence of the  $\Delta\varepsilon_\alpha$  of  $\alpha$ -relaxation for PLLA, PDLA and sc PLA.



**Figure 4. 12.** Temperature dependence of the dielectric strength  $\Delta\epsilon_{\alpha}$  of  $\alpha$ -relaxation for amorphous PLLA, PDLA and sc PLA.

In general, the dielectric relaxation strength  $\Delta\epsilon_{\alpha}$  decreases with temperature increasing [31, 37, 38]. Such a temperature dependence of  $\Delta\epsilon_{\alpha}$  was also reported in literature for other kinds of glass-forming systems, i.e. polymers and thin polymer films [31, 38, 39]. The obtained results indicate the same behavior of the dielectric strength  $\Delta\epsilon_{\alpha}$  for all the samples (Fig 4.12). In addition, the dielectric strength  $\Delta\epsilon_{\alpha}$  of sc PLA is close to the  $\Delta\epsilon_{\alpha}$  values of homopolymers. The maximum values of  $\Delta\epsilon_{\alpha}$  were reported at the lowest measurement temperature for each sample. The maximum of  $\Delta\epsilon_{\alpha}$  value ( $2.0 \pm 0.5$ ) is close to the previously reported results of Pluta et al. [40] for PLA ( $\sim 2.5$ ). The inset in Fig 4.12 presents the normalized  $\Delta\epsilon_{\alpha}$  by temperature ( $T\Delta\epsilon_{\alpha}$  normalized to the maximum value at lowest measured temperature) as a function of temperature for all the samples. Inset plot shows that  $T\Delta\epsilon_{\alpha}$  decreases with increasing temperature for each studied sample. As reported by Schönhalz [41], such temperature dependence caused by an increasing influence of intermolecular interactions between dipoles with decreasing temperature. Therefore, the stronger the temperature dependence of  $\Delta\epsilon_{\alpha}$ , the stronger the intermolecular interactions with decreasing temperature. Thus, such characteristics of the  $\alpha$ -relaxation can be defined by cooperativity character of the fundamental molecular motion responsible for the molecular dynamics. The size of the

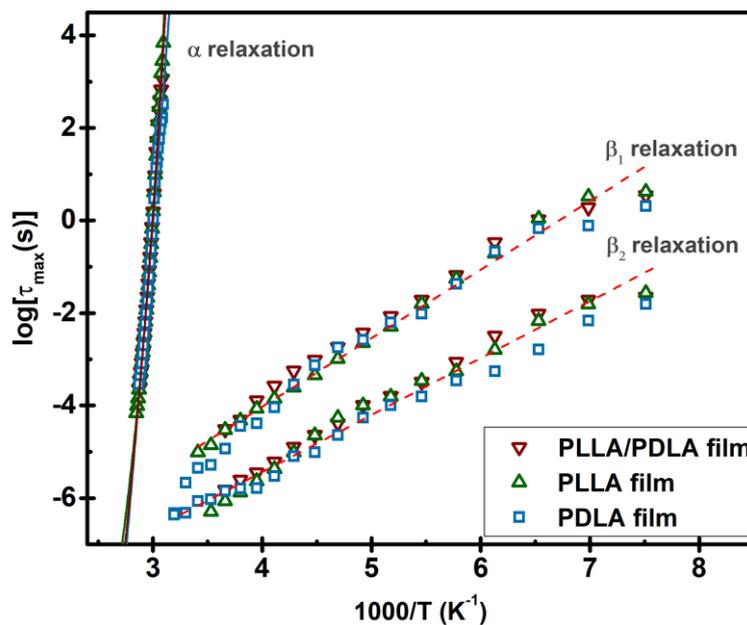
cooperative rearranging region increases with decreasing the temperature and, therefore, effective dipole moment increases [41].

#### 4.3.1.3 Relaxation map for $\alpha$ - and $\beta$ -relaxation

The relaxation map in Fig 4.13 shows the relaxation times  $\tau_{max}$  as a function of the inverse temperature for both  $\alpha$  and  $\beta$  processes. The temperature dependence of the relaxation time, i.e. the recorded  $\tau_{max}$  values, could be fitted and well described by the Arrhenius law as follows:

$$\tau = \tau_{0,A} \exp\left(\frac{E_a}{RT}\right) \quad (4.13)$$

where  $E_a$  is the activation energy of the  $\beta_1$  relaxation,  $R$  is the gas constant, and  $\tau_{0,A}$  is a pre-exponential factor.



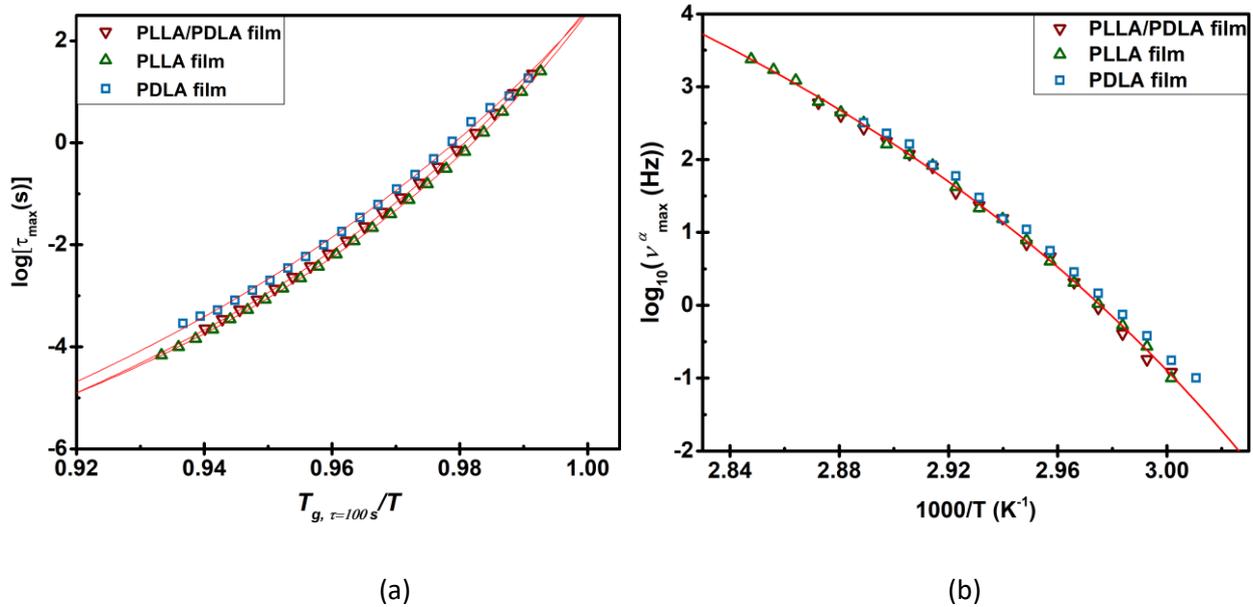
**Figure 4. 13.** Temperature dependence of structural  $\alpha$  and  $\beta$  relaxation times as a function of the inverse of temperature for amorphous PLLA, PDLA and sc PLA films.

The values of activation energy  $E_a$  are calculated as  $64 \pm 10$  kJ/mol for PLLA,  $57 \pm 12$  kJ/mol for PDLA, and  $65 \pm 10$  kJ/mol for sc PLA. Such results are in good agreement with the results for the other polyesters –  $79 \pm 10$  kJ/mol for PET [42] and 36 - 46 kJ/mol for PLA [28].

The values of  $E_a$  correspond to the local motions of the polar subunits of all the samples. In addition, the  $E_a$  value obtained for sc PLA is the same that the value obtained for PLLA and PDLA. Such results can be at the origin of the similarity in the chemical structure of the repeating units of homopolymers and sc PLA (Fig 2.1).

#### 4.3.1.4 Angell's and Stickel's plot of $\alpha$ relaxation

To investigate molecular dynamics in  $\alpha$  relaxation process, Angell's and Arrhenius plots are studied and presented in Fig 4.14. The symbols correspond to the experimental data and the red lines represent the VFT fits.  $T_g$  corresponds to  $\tau = 100$  s and  $\nu_{max}^\alpha$  is a relaxation rate, which is equivalent to  $1/\tau_{max}$ , where  $\tau_{max} = 1/2\pi f_{max}$  [43].



**Figure 4. 14.** (a) Relaxation time  $\tau_{max}$  as a function of normalized temperature  $T_{g, \tau=100s}/T$  (the Angell's plot), (b) Arrhenius plot of the amorphous PLLA, PDLA and sc PLA films for the  $\alpha$  relaxation process.

The temperature dependence of the relaxation time of the  $\alpha$ -relaxation can be fitted by a Vogel-Fulcher-Tamman law (VFT) by the following equation:

$$\tau_{max} = \tau_0 \exp\left(\frac{DT_0}{T-T_0}\right) \quad (4.14)$$

where  $\tau_{max}$  is the relaxation time at the maximum of the  $\alpha$  relaxation,  $D$  is the dimensionless parameter defined as the steepness strength,  $T_0$  is a reference temperature - so called Vogel

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POLYLACTIDE**

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temperature, and  $\tau_0$  is a pre-exponential factor. The Angell's plots in Fig 4.14 are all practically superimposed. Furthermore, the temperature dependence of the relaxation frequency corresponding to the dielectric  $\alpha$  relaxation peak  $\nu_{max}^\alpha$  is superimposed. Such a similarity can be explained by the similar dynamic glass transition temperature as shown in Table 4.2. The dielectric value of the glass transition temperature is usually selected as a temperature at a relaxation time equal to 100 s. The temperatures obtained by DRS at relaxation time equal 10 s match better with the calorimetric glass transition temperatures obtained by MT-DSC with a period of 60 s [44]. In literature, there are several studies of dielectric values of the glass transition temperature showing good deal with the values observed by thermal techniques, such as DSC and MT-DSC [45, 46]. For better fit of relaxation data in the relaxation temperature range, a second VFT law is required to define the temperature dependence of the relaxation time depending on the material [47, 48].

Samples	$T_g(\tau=100 \text{ s})$ (K)	$T_g(\tau=10 \text{ s})$ (K)	$T_g$ (K)	$D$	$T_0$ (K)	$\log(\tau_0)$ (s)	$E_\alpha$ (kJ mol <sup>-1</sup> )
PLLA	328	331	332	3.5	296	-13	64.4
PDLA	328	330	331	5.7	299	-15	57.3
Sc PLA	329	332	332	4.3	299	-14	64.7
PLA [49]	327	327		5	292	-14	

**Table 4. 2.** Results obtained by DRS: the glass transition temperature at  $\tau= 100$  and 10 s; dimensionless parameter of the VFT equation  $D$ ; reference temperature  $T_0$ ; the relaxation time extrapolated at an infinite temperature  $\tau_0$ ; the activation energy of the  $\beta_1$ -relaxation  $E_\alpha$ . The values of  $T_g$  are obtained by MT-DSC measurements.

The temperature dependence of the relaxation time of  $\alpha$ -process is investigated by analyzing the derivative method as proposed by Stickel et al. [50]. Stickel et al. presented the VFT description in glassy liquids which is associated with the linearity of  $\Phi_T$  as a function of  $T$ :

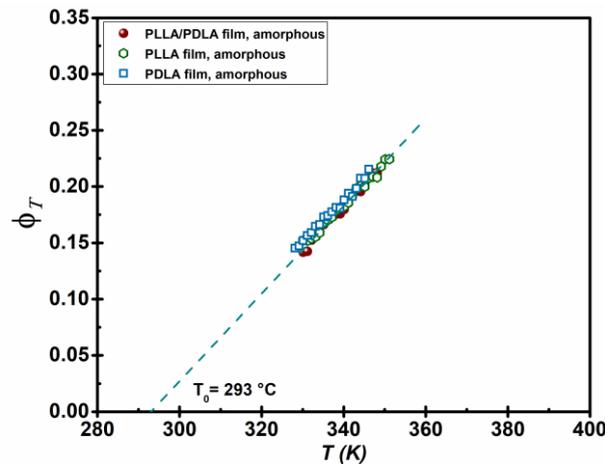
$$\Phi_T = \left[ \left( \frac{d \log_{10} \tau_{max}}{d(1000/T)} \right) \right]^{-1/2} \quad (4.15)$$

Stickel's plot is important key tool to determine the dynamic crossover temperature between two dynamic domains [47, 51]. Fig 4.15 depicts the derivative analysis of the dielectric relaxation time for all the samples. As shown, a single linear fit is sufficient to examine the behavior on the entire temperature range. For quantitative analysis, the VFT parameters  $B$  ( $B$

$= D_T \times T_0$ ) and the temperature  $T_0$  ( $\Phi_T = 0$ ) were estimated from derivative-based analysis by linear regression as follows [38]:

$$\left[ \left( \frac{d \log_{10} 1/\tau_{max}}{dT} \right) \right]^{-1/2} = \left( \frac{D_T}{2.303} \right)^{-1/2} (T - T_0) \quad (4.16)$$

The value of  $T_0$  and  $D_T$  are determined to be  $293 \pm 5$  K and  $4.12 \pm 1$  (as an average for three samples), respectively, so the parameter  $B$  is calculated to be  $1207 \pm 100$  K. The parameters obtained from the Stickel's analysis have good agreement and are similar for all the samples. This result is in good agreement with previous work on neat PLA presented by Rijal et al. [49]. All these parameters correspond to the fragility index values of the glass-forming liquids. By taking into account VFT fits and the derivative-based (Stickel's) analysis, uncertainty for the parameter  $B$  can be taken in the range of  $\pm 100$ , so Stickel's analysis can be also used to export VFT parameter for fragility investigation.



**Figure 4. 15.** The derivative-based analysis of the temperature evolution of dielectric relaxation data, so called Stickel's plot. Light blue dash line corresponds to the linear fit of the relaxation data.  $T_0$  is the Vogel temperature.

### 4.3.2 The fragility index $m$ by DRS

The next step of analysis for glassy polymers is calculating the fragility index  $m$  to extend quantitative comparison. High values of the fragility index  $m$  demonstrate "fragile" materials with a non-Arrhenius temperature dependence of the relaxation time  $\tau_\alpha$  approaches to  $T_g$ . The systems called "strong" supercooled liquids represent linear Arrhenius-like

temperature dependence of the relaxation time on the Arrhenius diagram. As defined, for  $m > 30$  liquids are considered as “fragile”, whereas for  $m < 30$  they are defined as “strong”. The degree of deviation from Arrhenius-type temperature dependence near  $T_g$  allows a classification of glass-formers on the basis of the structural relaxation, so called “fragility concept” by Angell et al. [52, 53] as follows:

$$m = \left. \frac{d \log(\tau_{max})}{d\left(\frac{T_g}{T}\right)} \right|_{T=T_g} \quad (4.17)$$

where  $T_g / T$  is the temperature normalized with respect to the glass transition temperature  $T_g$ . All parameters which are necessary to calculate the fragility index are gathered in Table 4.2.

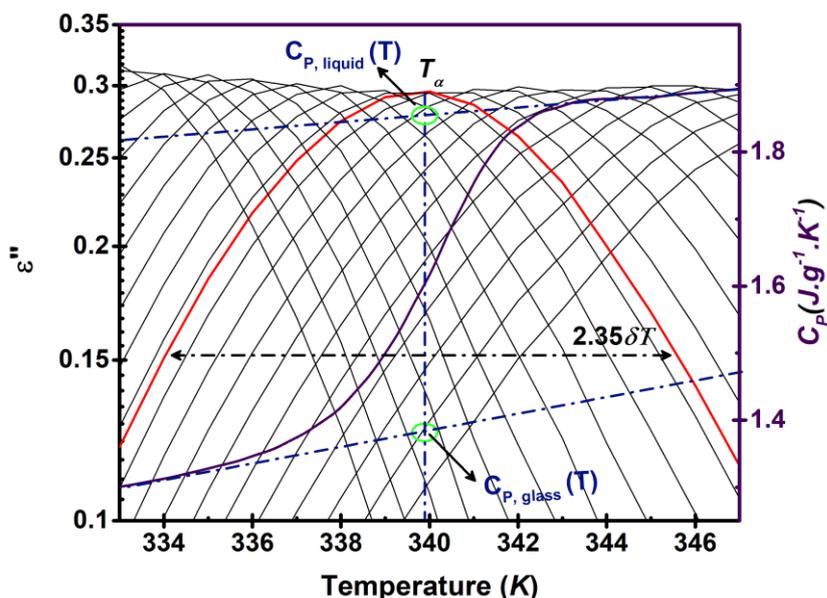
The calculated values of  $m$  for homopolymers and sc PLA are 146 and 150, respectively. Therefore, such results correspond to “fragile” glass-formers. Such behavior depends on the molecular structure with extended van der Waals interactions and/or hydrogen bonds between polymer chains [52, 54]. In previous works, the fragility index  $m$  of PLAs was reported as 144 [49], 155 [55], 149 [56]. According to the accuracy of  $\pm 10\%$  for the fragility index calculation, we can assume that amorphous sc PLA has practically the same fragility index as neat PLA and homopolymers. Furthermore, the values of the fragility index  $m$  determined from DRS analysis match well with the values of fragility index  $m$  calculated from FSC measurements by VFT fitting.

### 4.3.3 Molecular mobility and dynamic heterogeneity

Previously, Schick et al. [57] combined DRS, MT-DSC and Ac-chip calorimetry in order to study molecular mobility of different polymers, such as polystyrene (PS) and poly(methyl methacrylate) (PMMA). They found that the different experimental techniques show consistent results of cooperativity length obviously correlated with the structural relaxation temperature and time dependence.

**4.3.3.1 Method of extraction the parameters for cooperativity size calculation**

Typical length scale of cooperativity ranging from 1 to 3.5 nm was provided at the glass transition temperature for different glassy polymers. Besides, the relationship between the cooperativity, the glass transition temperature and the fragility index has been proposed [58]. It is found that the CRR size increases with increasing fragility. Fig 4.16 represents isochronal spectra of the dielectric loss in the frequency range of  $2.10^6$  Hz down to 0.1 Hz combined with the heat capacity curves from MT-DSC as a function of the temperature for sc PLA (same calculation was applied for all the amorphous samples). The dielectric spectra were plotted by subtracting the contribution of the conductivity and the secondary  $\beta$  relaxation. The observed relaxation peak corresponds to the segmental ( $\alpha$ -) relaxation process. The CRR length can be estimated by using the temperature fluctuation of the amorphous phase by Eq 1.20 as Donth proposed [59]. All the quantities used in Donth's approach [59] were determined from DRS and MT-DSC investigation.



**Figure 4. 16.** Dielectric loss (without conductivity) from DRS measurements and heat capacity from MT-DSC measurement as a function of temperature. The blue dash lines present the heat capacity in the glassy and liquid states. The red line corresponds to the Gaussian fit of a dielectric loss spectra.

As shown in Fig 4.16, the temperature fluctuation  $\delta T$  (related to the standard deviation  $\sigma_T$  of the Gaussian peak function) is calculated as  $\delta T = \text{FWMH} / 2.35$  and the dynamic glass

transition temperature  $T_\alpha$  (corresponds to the maximum of the Gaussian peak) is exported from Gaussian fit of the imaginary part of MT-DSC and DRS spectra by following equation:

$$C''(T) \text{ or } \varepsilon''(T) = \frac{A}{\sigma_T \sqrt{2\pi}} \exp\left(-\frac{(T-T_\alpha)^2}{\sigma_T^2}\right) \quad (4.18)$$

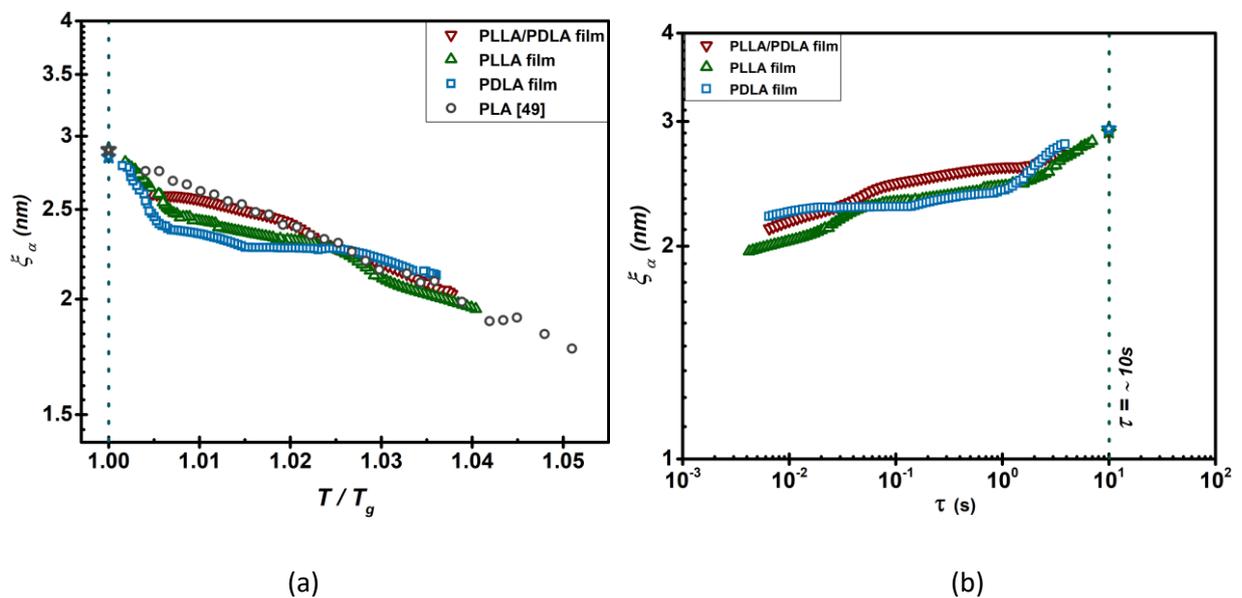
where  $A$  corresponds to the peak area,  $\sigma_T$  is the standard deviation and  $T_\alpha$  is the dynamic glass transition temperature. The value  $\Delta C_p^{-1}$  is the difference in the inverse of the isobaric heat capacity between the glass and the liquid at  $T_\alpha$  and exported from the calorimetric investigation, as shown in Fig 4.16.

Therefore, combination of MT-DSC and DRS allows us to calculate the temperature dependence of the cooperativity length  $\xi_\alpha$  (by Eq 1.20) and the cooperativity number of relaxing entities  $N_\alpha$  (by Eq 1.21) as well as the number of dynamically correlated units  $N_c$  (by Eq 1.15) in a wide temperature and relaxation time range from the onset of cooperativity in the crossover region to the calorimetric glass transition temperature  $T_g$ .

#### 4.3.3.2 The cooperativity length $\xi_\alpha$ and cooperativity degree $N_\alpha$

The cooperativity length  $\xi_\alpha$  and the cooperativity degree  $N_\alpha$  were estimated in a wide range of temperature and relaxation time for each investigated sample. The cooperativity length  $\xi_\alpha$  associated with the  $\alpha$ -relaxation as a function of temperature and relaxation time determined from MT-DSC and DRS measurements is presented in Fig 4.17. The results shows that the cooperativity length  $\xi_\alpha$  depends on the relaxation time and the glass transition temperature. In addition, the cooperativity degree  $N_\alpha$  is presented in Fig 4.18 to investigate the temperature dependence of  $N_\alpha$  and compare it with other systems. A non-linear increase of the cooperativity length  $\xi_\alpha$  and the cooperativity degree  $N_\alpha$  with decreasing temperature and increasing relaxation time is observed for each sample. All studied samples have similar behavior regarding temperature dependence of the cooperativity parameters. In addition, the similarity in the cooperativity length of homopolymers and sc PLA could also be correlated with the results of similar fragility index  $m$  of samples. It should be noted that the extrapolation of the cooperativity length  $\xi_\alpha$  estimated from DRS over a wide range of relaxation time and temperature fits very well with the values estimated by MT-DSC (star

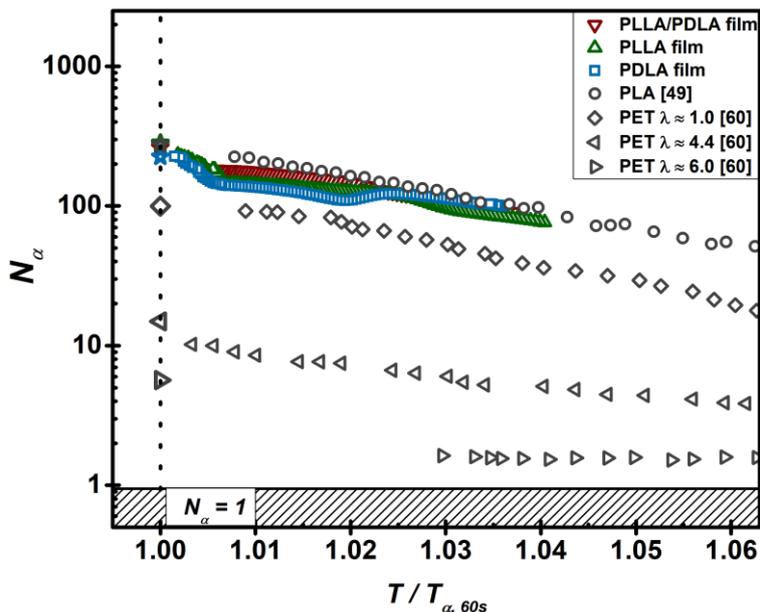
shape symbols) at calorimetric glass transition  $T_\alpha$  at  $\tau \approx 10$  s as well as the cooperativity number  $N_\alpha$ . However, such a non-linear temperature dependence of  $\xi_\alpha$  and  $N_\alpha$  can be related to the temperature dependence of the relaxation time. The cooperativity length  $\xi_\alpha$  at the calorimetric  $T_g$  for all the samples is quite similar, i.e.  $2.9 \pm 0.2$  nm for PLLA and sc PLA, and  $2.8 \pm 0.2$  nm for PDLA, independently from the intermolecular interactions obtained through stereocomplexation.



**Figure 4. 17.** The cooperativity length  $\xi_\alpha$  as a function of (a) normalized temperature at  $T_g$  and (b) relaxation time for each sample obtained from DRS measurements by estimated the calorimetric glass transition temperature at period 60 s,  $\tau \sim 10$  s. The star shaped symbols at  $T/T_g = 1$  correspond to the results derived from MT-DSC analysis at  $T_\alpha$  at  $\tau \approx 10$  s.

Furthermore, as depicted in Fig 4.17, the values are rather similar to the results obtained by Rijal et al. [49] for commercial PLA. It is interesting to compare the results of our systems with the results of Hamonic et al. [60] obtained for another polyester, PET with different microstructure, i.e. wholly amorphous and isotropic PET, and semi-crystalline PET drawn at two different draw ratio ( $\lambda$ ) (Fig 4.18). Hamonic et al. have shown a huge impact of the microstructure on the temperature dependence of the cooperativity.

As a consequence, if the amorphous phase is anisotropic and constraint by crystallites (i.e. highly drawn PET [60]), the CRR size and  $N_\alpha$  are drastically smaller. These results reveal that cooperativity could vary a lot in the amorphous part of the polymer, but in the case of amorphous PLA, stereocomplexation has no significant influence.

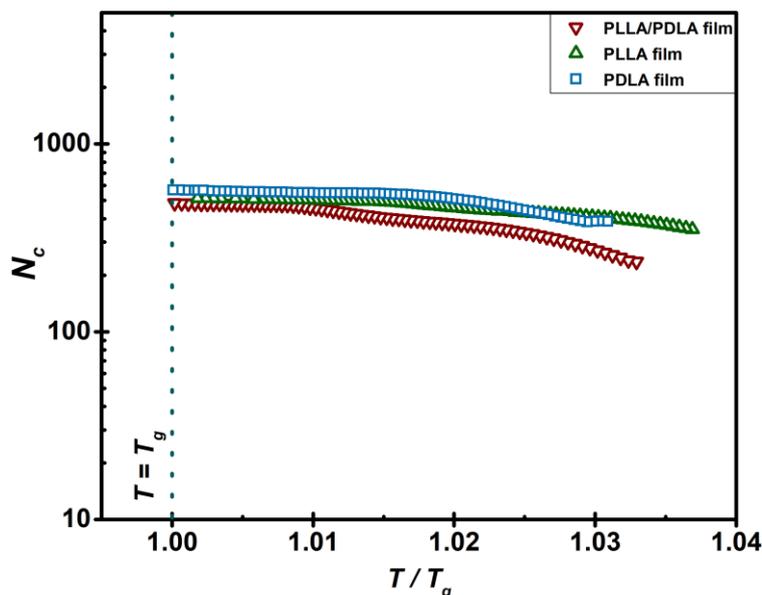


**Figure 4. 18.** The cooperativity degree  $N_\alpha$  as a function of normalized temperature. The star shaped symbols at  $T/T_{\alpha, 60s} = 1$  correspond to the results obtained from MT-DSC analysis at  $T_\alpha$  at  $\tau \approx 10$  s.

#### 4.3.3.3 Dynamic heterogeneity and comparison of $N_C$ and $N_\alpha$

The number of dynamically correlated segments  $N_C$  of all the samples is calculated by Eq 1.15 and is depicted in Fig 4.19 as a function of temperature normalized to  $T_g$ . The calculated values of  $N_C$  for each sample exhibit similar temperature dependence. The slight increase of the number of dynamically correlated segments  $N_C$  with decreasing temperature towards the glass transition temperature is observed in Fig 4.19. These results show good relevance with the tendency for all the amorphous samples. Furthermore, the increase of  $N_C$  with decreasing temperature is independent from the chemical structure of the glass-forming liquid samples. Therefore, such increase can be explained by two regimes: the first fast regime is observed at short relaxation times and the second one, much slower, is noted for relaxation times towards the glass transition. Thus, the extent of spatial correlation in the molecular dynamics so grows as temperature approaches the glass transition. This result is supported by

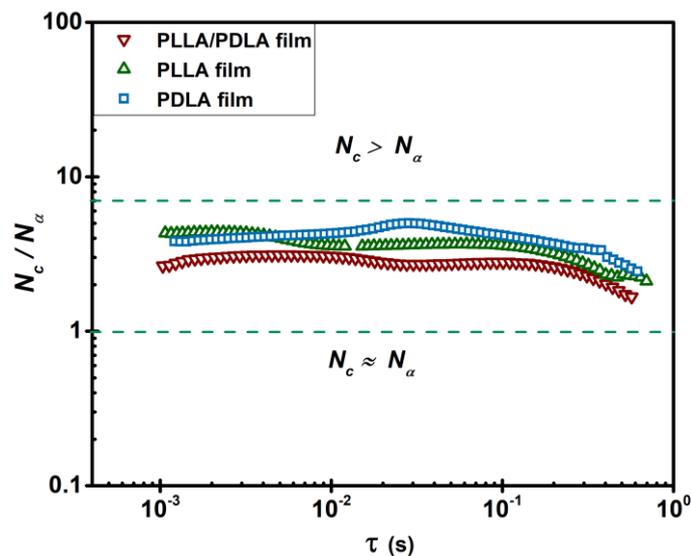
several works [49, 61, 62] and can be predicted also by the mode coupling theory [63]. These results support the common evolution of  $N_C$  during the relaxation for a large family of glass-forming liquids.



**Figure 4. 19.** The number of dynamically correlated segments  $N_C$  as a function of temperature normalized at  $T_g$  for all the samples.

Although the direct comparison of  $N_C$  and  $N_\alpha$  is insignificant due to the shift in absolute values, similar behavior can be observed for all the samples. The increase of  $N_\alpha$  with increasing relaxation time can be also described by a short relaxation time, a single power law and a long relaxation times up to  $T_g$  ( $\tau = 100$  s), as the variations of  $N_C$ . In order to compare more precisely the number of dynamically correlated segments  $N_C$  with the cooperativity degree  $N_\alpha$  calculated from the Donth's approach, the ratio  $N_C / N_\alpha$  is presented in Fig 4.20 as a function of the relaxation time. When close to the  $T_g$  (in the range of  $\tau_{Tg} \approx 10$  s – 100 s), the values of  $N_C$  and  $N_\alpha$  are rather similar ( $N_C / N_\alpha \sim 1$ ). However, these values start to deviate with the relaxation time decrease. Therefore, we may suppose that such similarities can be found in the physical definition of  $N_\alpha$  and  $N_C$ . The number of dynamically correlated segments  $N_C$  is calculated based on the 4-point correlation approach, different from the Donth's approach, since it allows us to calculate the number of molecules whose dynamics are correlated depending on the temperature fluctuations. However, with a relaxation time decrease, the

ratio of  $N_c / N_\alpha$  slightly increases and remains constant. For these two approaches, the temperature or the relaxation time associated with the value  $N_c$  or  $N_\alpha = 1$  corresponds to the limit of the cooperativity, and defines a “cross-over” point in Arrhenius diagram. From the results of  $N_c$  and  $N_\alpha$  as a function of relaxation time, the “cross-over” point is observed at the relaxation time  $\tau \sim 10^{-13}$  s depending on the  $N_c = 1$ , but it is at the range of  $\tau \sim 10^{-11}$  s according to  $N_\alpha = 1$  for all the samples (as an average of three samples). Results show that  $N_\alpha$  decreases faster than  $N_c$  with a decreasing relaxation time. According to the relaxation time investigation, the different asymptotic behaviors are observed in these two models. These new and original results of sc PLA compared to the homopolymers indicate that the origin of the cooperative molecular motions for both  $N_c$  and  $N_\alpha$  of sc PLA is independent from stereocomplexation. Therefore, the different values of the relaxation time and the different definitions between correlated and cooperative motions (especially for very short relaxation times) resulting in the correlated motions appear at a time scale lower than the cooperative motions.



**Figure 4. 20.** The ratio of  $N_c/N_\alpha$  as a function of the relaxation time.

## **Conclusion**

In this chapter, the influence of the tacticity on the molecular dynamic of poly(lactides) has been investigated in terms of the segmental ( $\alpha$ -process) relaxation by FSC and DRS measurements. It is found that fictive temperatures and glass transition values determined by different techniques (by FSC, DSC, MT-DSC or DRS) have same values for PLLA, PDLA and sc PLA. The kinetics of enthalpy recovery of amorphous sc PLA and homopolymers (PLLA and PDLA) was investigated by FSC measurements. The results show that the enthalpy recovery has a single step decay towards equilibrium and glasses are allowed to completely relax in the range of aging time. It is found that the relaxation kinetics of the glass towards the enthalpy recovery of the supercooled liquid is independent on the tacticity in the stereocomplex structure.

Furthermore, the structural relaxation investigation was carried out by DRS analysis for wholly amorphous samples. The  $\alpha$ - and  $\beta$ -processes of wholly amorphous samples were examined to observe the dielectric properties such as relaxation time, dielectric strength, and fragility. All the samples have the similar dielectric properties independently on the tacticity existence. The cooperativity length  $\xi_\alpha$  estimated from Donth's approach for the all samples was studied and found to be very similar at the glass transition ( $\approx 2.9$  nm) and in a wide temperature range. It is shown that the cooperativity length  $\xi_\alpha$  has no dependence on the chemical structure, while the cooperativity number  $N_\alpha$  has the strong dependence on the microstructure. Moreover, no effect of stereocomplexation on the origin of the cooperative molecular motions is observed. The correlated motions appear at a time scale lower than the cooperative motions for all the samples. It is found that the kinetics of the segmental relaxation of wholly amorphous sc PLA and the amorphous phases of PLAs are the same. In addition, the results of  $\beta$ -process testifies that the local movements of polymer chain are independent from the stereocomplexation impacts on the local interactions in case of amorphous samples, although stereocomplexation led to generate different crystalline phase.

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**CHAPTER 5. MICROSTRUCTURE INFLUENCE ON MECHANICAL AND BARRIER PROPERTIES OF STEREOCOMPLEX PLA**

The influence of the stereocomplexation on the barrier properties of PLA-based materials towards water and gases (N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>) is discussed in the present chapter. Special attention is paid on the water permeability as a function of temperature. Transport properties towards water were investigated for both the amorphous and semicrystalline samples. In addition, gas permeation measurements were carried out to investigate the barrier properties of semicrystalline samples. The transport properties were correlated with the microstructure investigations.

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### 5.1 Microstructure investigations

DSC and XRD measurements were carried out to study the microstructure of the crystalline phases of polymer films. As depicted in Fig 5.1, the sc PLA crystal has higher melting temperature (about 50 °C) than  $\alpha$  crystals.

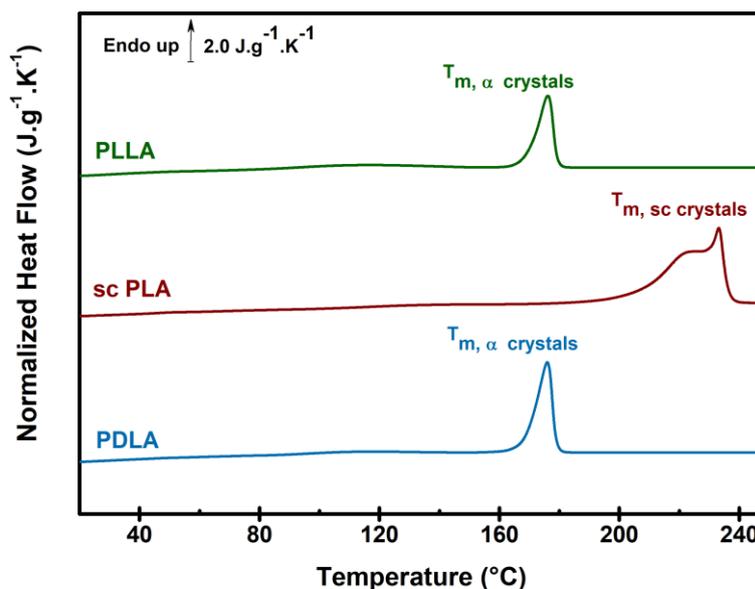


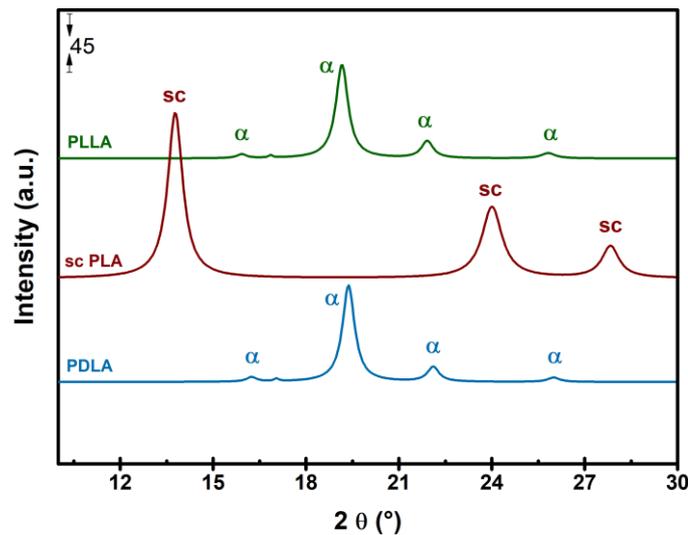
Figure 5. 1. Normalized heat flow curves of PLLA, PDLA, and sc PLA films.

The values of thermal characteristics (i.e. glass transition temperature  $T_g$ , heat capacity step  $\Delta C_p$ , crystallinity degree  $X_c$ ) calculated from DSC measurements are presented in Table 5.1 for homopolymers and sc PLA. The  $T_g$  values of semicrystalline PLAs are higher than those of amorphous PLAs due to the strong coupling between the phases caused by the low crystallization temperature (Table 5.1) (it is also discussed previously in Chap. 3 (Fig 3.23)). The  $X_c$  value of sc PLA is slightly higher than that of homopolymers. Also, semicrystalline sc PLA has smaller heat capacity step which is closed to the limit of detection as compared to homopolymers. Therefore, we observed broad glass transition range. The crystalline structure was also confirmed by XRD analysis as presented in Fig 5.2. The presence of only one crystalline phase was revealed by the reflection peaks at  $2\theta = 19^\circ$  and  $22^\circ$  for  $\alpha$  crystals and  $2\theta = 14^\circ$ ,  $24^\circ$  and  $28^\circ$  for sc crystals.

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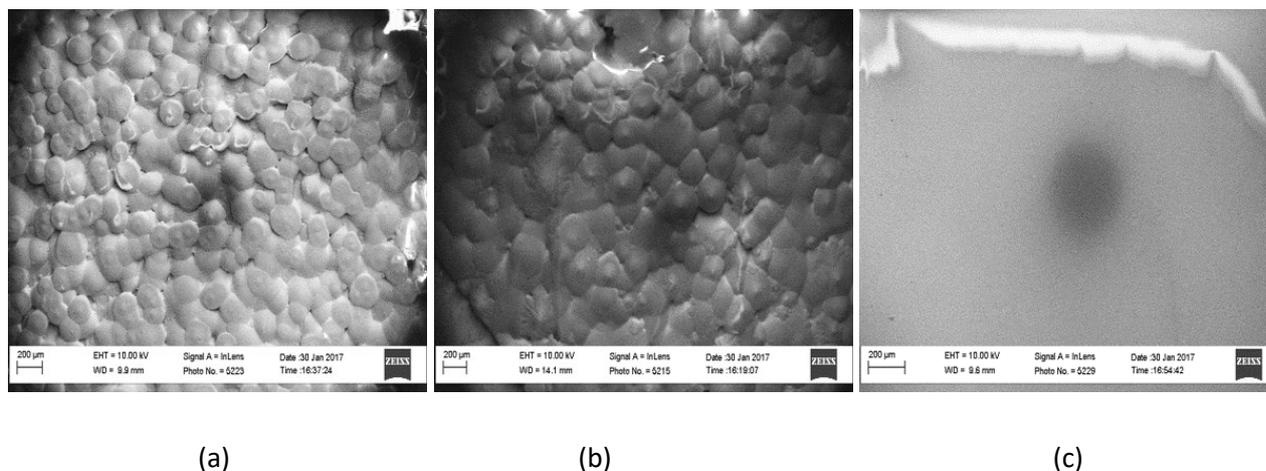
Samples	$T_g$ (°C)		$\Delta C_p$ (J.g <sup>-1</sup> .K <sup>-1</sup> )		3-phases of semicrystalline samples		
	amorphous	semicrystalline	amorphous	semicrystalline	$X_c$ (%)	$X_{MAP}$ (%)	$X_{RAF}$ (%)
PLLA	59	75	0.46	0.16	48	35	17
PDLA	61	76	0.46	0.19	48	41	11
Sc PLA	59	69	0.54	0.12	54	22	24

**Table 5. 1.** Thermal properties of studied amorphous and semicrystalline samples.



**Figure 5. 2.** XRD spectra of PLLA, PDLA and sc PLA films.

In addition, surface morphology of PLLA, PDLA and sc PLA films was examined by SEM (Fig 5.3). As one can see from Fig 5.3, homopolymers films are characterized by a granular structure (Fig 5.3a and b) while sc PLA film has a smooth compact structure (Fig 5.3c). Besides, no porosity is observed for all film samples. Such difference in surface morphology can be caused by the formation of the different crystallites types (i.e. pseudoorthorhombic which contains  $10_3$  helices for homo-crystals and triclinic in which *L*- and *D*-lactide are packed as parallel taking  $3_1$  helical conformation for sc crystals [1]) during the crystallization through solvent evaporation.



**Figure 5. 3.** SEM images of (a) PLLA, (b) PDLA, and (c) sc PLA film.

## 5.2 Barrier Properties

Several works can be already found in the literature dealing with the improvement of the barrier properties of PLA-based membrane in order to fulfill the industrial packaging requirements. Tenn et al. [2] found that the incorporation of nanoparticles (Cloisite 30B (C30B)) in the PLA matrix decreased the water permeability with the increase of the nanoclay content. It is also shown that the transport properties depend strongly on the dispersion state of the nanoclay platelets and their orientation in the PLA matrix. Tenn et al. [3] also found that despite the fact that the surface modification of PLA films by hydrophobic plasma treatments improved the water barrier properties, no influence was observed on the transport properties towards oxygen. Besides the nanocharges [4-6], the transport properties can be improved by modifying the morphology of the polymer films [7-10]. For example, Messin et al. [11] revealed that the use of semicrystalline poly(butylene succinate-co-butylene adipate (PBSA) polymer improved the barrier properties of PLA in the case of multilayer PLA/PBSA film. In addition, Delpouve et al. [12] confirmed that the presence of  $\alpha'$  crystals had no influence on the PLA water permeability, but the drawing processes allowed improving the PLA barrier properties due to the crystals shape and increase in crystallinity degree. Therefore, it was shown that the crystalline and amorphous phase organizations are the main parameter of the barrier properties improvement in the case of PLA [12]. Furthermore, Bai et al. [13] reported

that the shape of crystals had influence on the barrier properties. The oxygen barrier performance of PLA was improved through constructing parallel-aligned shish-kebab-like crystals with well-interlocked boundaries.

In our work, the transport properties of PLA-based materials obtained by stereocomplexation are investigated and compared to those of homopolymers. The permeation measurements for homopolymers (PLLA, PDLA) and sc PLA are performed towards water and gases (N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>) and are correlated with the molecular mobility study (i.e. the CRR volume and size).

### **5.2.1 Water barrier properties**

The water permeation measurements were carried out at controlled temperature 25 °C, 45 °C and 65 °C by using a home-made apparatus (as detailed in chapter 2). The water permeation flux  $J$  was recorded at the dry interface of the investigated sample. To annihilate the influence of the film thickness, the water permeation kinetics curves were analyzed by using the reduced scale  $JL = f(t/L^2)$ . Each permeation curve may be divided into three parts [14]. At the beginning of the measurement no water flux is determined, i.e.  $JL = 0$ . Then, after a while, water molecules start to pass through the film and the strong increase of the water flux is observed. Such increase corresponds to the transient step, from which the diffusion coefficient  $D$  can be estimated. After transient step, the steady state (stationary state, during which the water flux remains constant  $J_{stat}$ ) is reached and the permeability coefficient  $P$  can be calculated as the amplitude of the curve. Therefore, the reduced water permeation flux was plotted as a function of the reduced time, and the values of the water permeability coefficient  $P$  were determined from the flux at stationary state  $J_{stat}$  by using the following equation:

$$P = \frac{J_{stat}L}{\Delta a} \quad (5.1)$$

where  $L$  is the thickness of the film sample and  $\Delta a$  is the activity difference between the two faces of the sample,  $\Delta a = 1$  in case of liquid water, i.e. liquid water on the upstream and dry

gas on the downstream face of the film. The permeability coefficient  $P$  is usually expressed in Barrer units as:

$$1 \text{ Barrer} = 10^{-10} \frac{\text{cm}^3 (\text{STP})\text{cm}}{\text{cm}^2 \text{ s cmHg}} \quad (5.2)$$

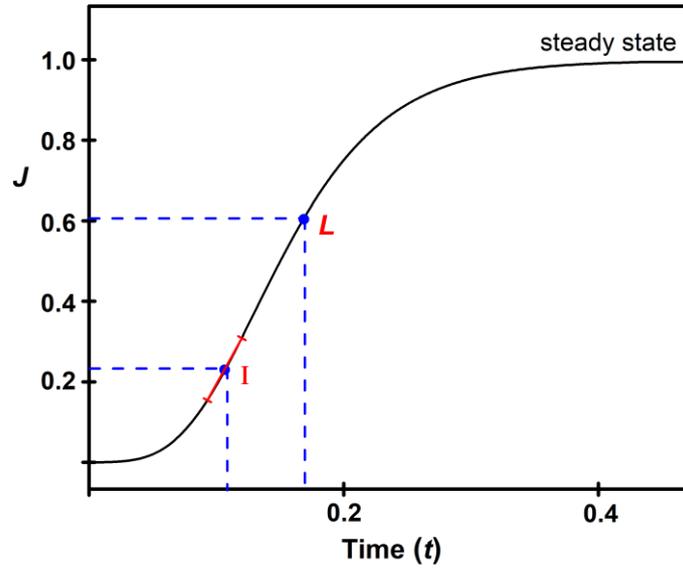
The materials often used such as epoxy resins or polyesters, may suffer from plasticization, local strain and chemical modifications/degradations over a long period of time [15]. In order to examine the possible plasticization effect induced by the water molecules in the studied samples, the dimensionless curve  $j = f(\tau)$  calculated from Fick's equations was defined by Follain et al [16], where  $j = \frac{J}{J_{stat}}$ ,  $\tau = \frac{D.t}{L^2}$  and  $D$  is the diffusion coefficient assumed as a constant value. The diffusion coefficient  $D$  was calculated from the transient regime of the water permeation flux according to the Fickian behavior. The two distinct diffusion coefficient noted as  $D_{0.24}$  and  $D_{0.62}$  can be distinguished:

$$D_{0.24} = \frac{L^2 \times 0.091}{t_{0.24}} \quad (5.3)$$

and

$$D_{0.62} = \frac{L^2}{t_{0.62}} \quad (5.4)$$

$D_{0.24}$  and  $D_{0.62}$  were calculated at two different times of the permeation process (assumed as  $t_I$  and  $t_L$ ) that corresponds to the inflection point I ( $j_I = 0.24$ ,  $\tau = 0.091$ ) and to the time-lag point L ( $j_L = 0.62$ ,  $\tau = 1/6$ ) of the theoretical flux curve, respectively, as detailed in Fig 5.4 [17].



**Figure 5. 4.** Representation of the flux ( $J$ ) as a function of time ( $t$ ) to determine the  $P$  and  $D$  coefficient.

The plasticization effect induced by water molecules maybe described by an exponential dependence of diffusion coefficient with water concentration by:

$$D = D_0 e^{\gamma \cdot C} \quad (5.5)$$

where  $D_0$  is the diffusion coefficient when the water concentration tends to 0,  $\gamma$  is the plasticization factor and  $C$  is the local permeant concentration in the membrane. Therefore, the maximum diffusion coefficient  $D_M$  can be determined as follows [15]:

$$D_M = D_0 e^{\gamma C_{eq}} \quad (5.6)$$

where  $C_{eq}$  is the water concentration within the studied sample at the stationary state. Therefore, the mean integral diffusion coefficient  $\langle D \rangle$  can be calculated as follows:

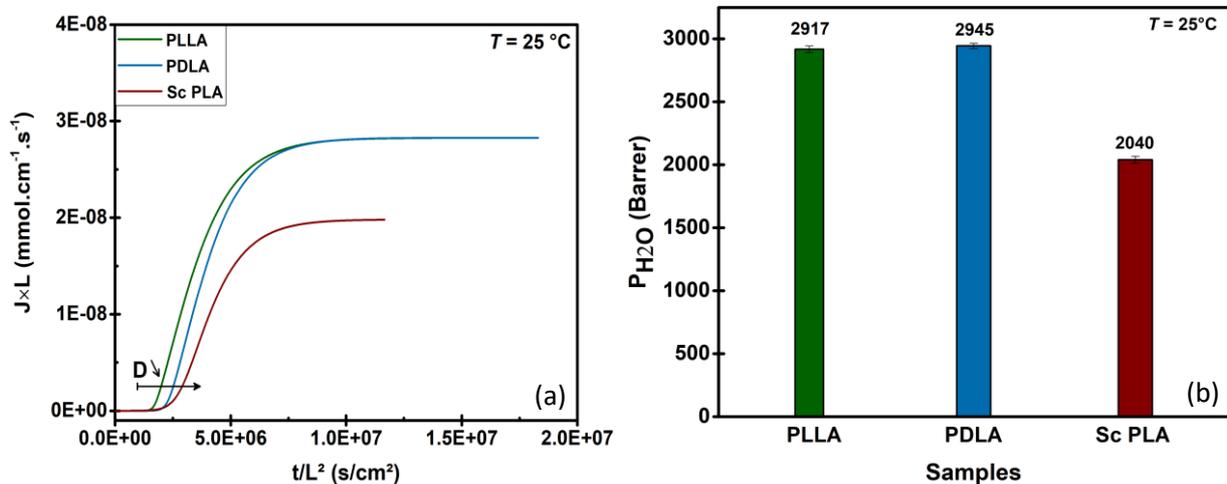
$$\langle D \rangle = \frac{1}{\Delta C} \int_{C_1}^{C_2} D(C) \cdot dC \quad (5.7)$$

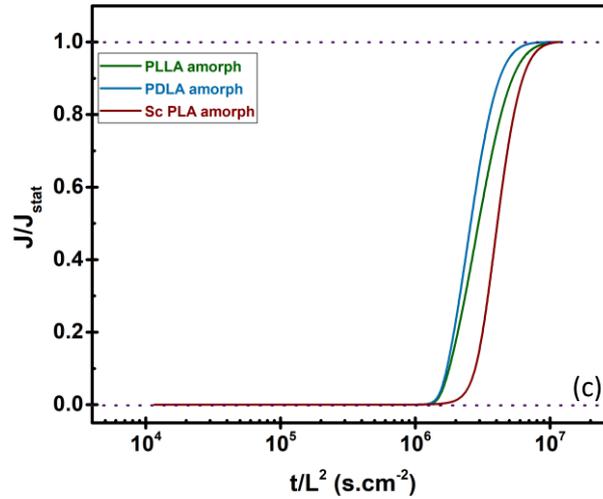
#### 5.2.1.1 Water permeability of amorphous samples

The water permeation tests of amorphous samples towards water were carried out at 25°C to examine the stereocomplexation impact on the barrier properties. The water permeation coefficient was calculated from the  $J^*L = f(t/L^2)$  curves (Fig 5.5a) and the values are presented in Fig 5.5b. Sc PLA reveals lower value of  $P$  as compared to homopolymers.

## CHAPTER 5. MICROSTRUCTURE INFLUENCE ON MECHANICAL AND BARRIER PROPERTIES OF STEREOCOMPLEX PLA

Therefore, one can say that stereocomplexation improves the water barrier properties due to the intermolecular stereoselective interactions; although, the glass transition properties ( $T_g$ , CRR volume and size) and molecular dynamics of amorphous samples are the same (presented in Chap. 4). As seen in Fig 5.5a and c, an increase in the delay time in diffusion on the normalized curve of flux for sc PLA is observed compared to the homopolymers. The calculated values of  $D_0$  ( $C=0$ ) and  $D_M$  ( $C=C_{eq}$ ) are shown in Table 5.2. As one can see, the increase of the  $D$  values with the water concentration and the plasticizing factor  $\gamma > 0$  confirm the existence of plasticizing effect. According to the values of water concentration ( $C_{eq}$ ) at steady state, sc PLA films are characterized by lower value of  $C_{eq}$  compared to homopolymers. It can be also noted the higher affinity between water molecules and homopolymers was revealed. This affinity can be at the origin of the differences in barrier properties of the studied samples taking into account that intermolecular stereoselective interactions limit the water concentration in the film in case of sc PLA.





**Figure 5. 5.** (a) Evolution of the reduced flux of water permeation ( $J^*L$ ) as a function of reduced time ( $t^*L^2$ ), (b) water permeability coefficient  $P$  and (c) normalized water permeation curves as a function of the reduced time for amorphous PLLA, PDLA and sc PLA.

Samples	$P$ (Barrer)	$D_0 \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$D_M \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$\gamma C_{eq}$	$\gamma$ ( $\text{cm}^3 \cdot \text{mmol}^{-1}$ )	$C_{eq}$ ( $\text{mmol} \cdot \text{cm}^{-3}$ )
PLLA	$2917 \pm 26$	$2.4 \pm 0.3$	$14.6 \pm 2.7$	$1.8 \pm 0.2$	$4.3 \pm 0.1$	$0.42 \pm 0.03$
PDLA	$2945 \pm 21$	$1.6 \pm 0.4$	$18.0 \pm 2.9$	$2.4 \pm 0.1$	$5.8 \pm 0.4$	$0.42 \pm 0.04$
Sc PLA	$2040 \pm 28$	$1.3 \pm 0.1$	$18.7 \pm 2.7$	$2.7 \pm 0.2$	$8.8 \pm 1.8$	$0.30 \pm 0.05$

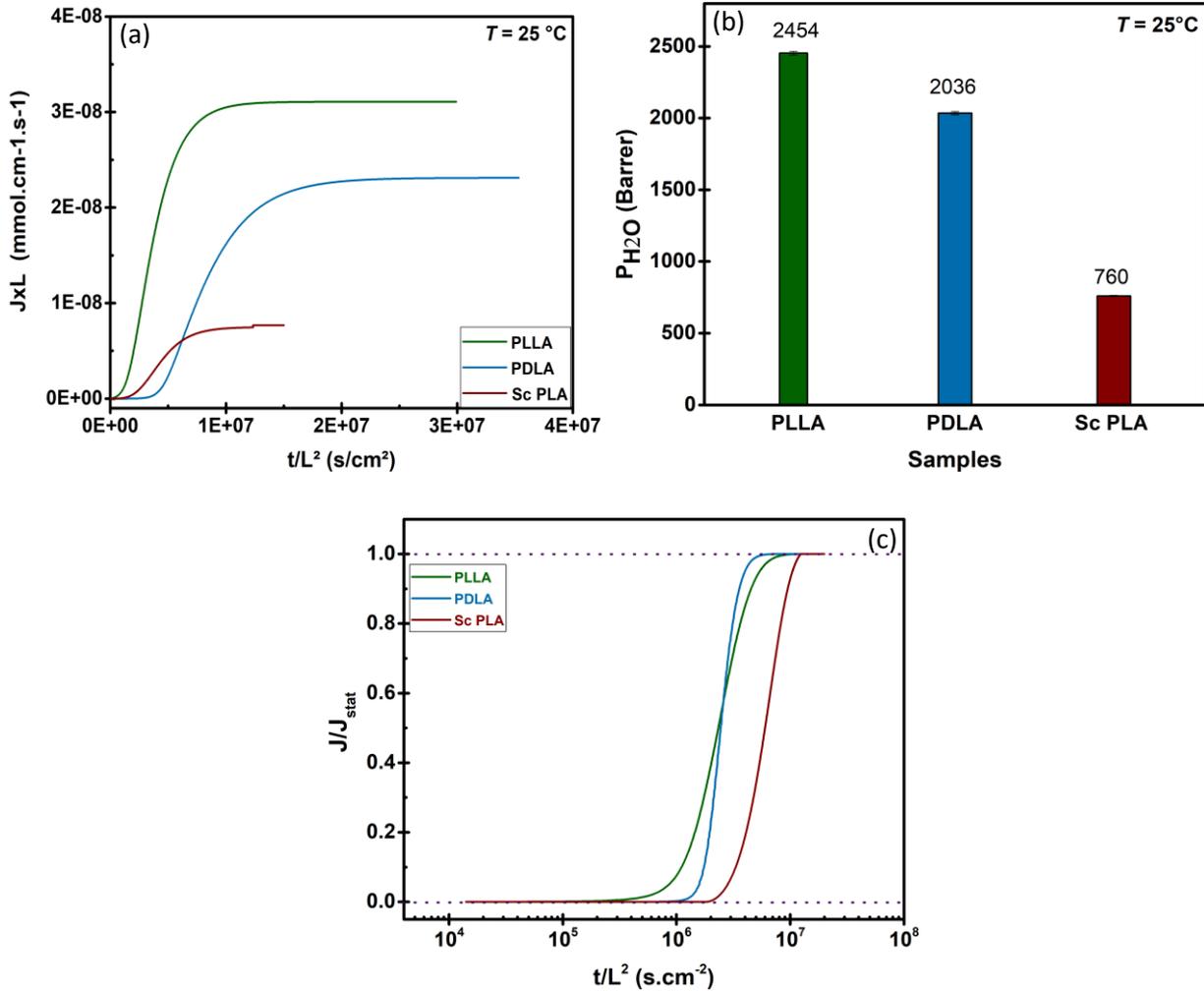
**Table 5. 2.** Water permeametric coefficients of amorphous PLLA, PDLA and sc PLA at  $25 \pm 1^\circ\text{C}$ .

### 5.3.1.2 Water permeability of semicrystalline samples

The water barrier properties of semicrystalline samples were studied at temperature of  $25^\circ\text{C}$ ,  $45^\circ\text{C}$  and  $65^\circ\text{C}$ . The results are presented for each measurements' temperature separately. According to the obtained results, the permeation flux through the sc PLA film is lower than that in the case of two homopolymers whatever the studied temperature. Such results obviously demonstrate an improvement of the barrier properties of PLA by the stereocomplexation and/or crystallization. Besides, it should be noted that while the permeability coefficient of sc PLA increases with  $T$ , the permeability of homopolymers decreases by increasing the temperature.

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- $T_{\text{permeation}} = 25^{\circ}\text{C}$



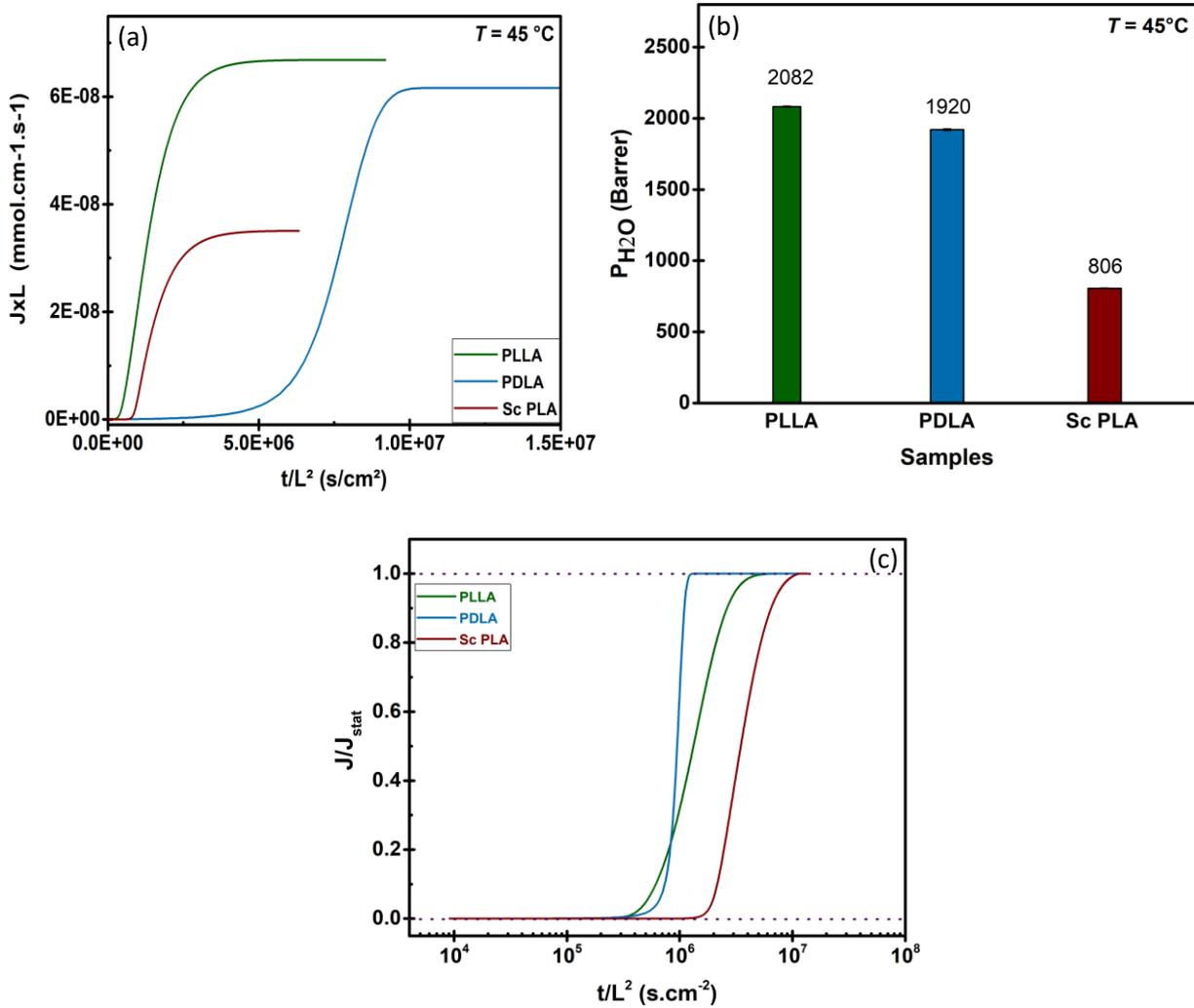
**Figure 5. 6.** (a) Evolution of the reduced flux of water permeation ( $J^*L$ ) as a function of reduced time ( $t^*L^2$ ), (b) water permeability coefficient  $P$ , and (c) normalized water permeation curves as a function of the reduced time for semicrystalline PLLA, PDLA and sc PLA at  $T_{\text{permeation}} = 25^{\circ}\text{C}$ .

25°C Samples	P (Barrer)	$D_0 \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$D_M \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$\gamma C_{\text{eq}}$	$\gamma$ ( $\text{cm}^3 \cdot \text{mmol}^{-1}$ )	$C_{\text{eq}}$ ( $\text{mmol} \cdot \text{cm}^{-3}$ )
PLLA	$2454 \pm 13$	$1.26 \pm 0.34$	$8.7 \pm 8.7$	$2.17 \pm 1.22$	$2.29 \pm 0.18$	$0.83 \pm 0.03$
PDLA	$2036 \pm 11$	$0.92 \pm 0.56$	$7.01 \pm 1.9$	$2.04 \pm 0.05$	$2.64 \pm 0.62$	$0.77 \pm 0.08$
Sc PLA	$760 \pm 5$	$1.76 \pm 0.05$	$4.74 \pm 1.7$	$0.99 \pm 0.02$	$4.00 \pm 1.23$	$0.25 \pm 0.02$

**Table 5. 3.** Water permeametric coefficients of semicrystalline PLLA, PDLA and sc PLA at  $25 \pm 1^{\circ}\text{C}$ .

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- $T_{\text{permeation}} = 45^{\circ}\text{C}$



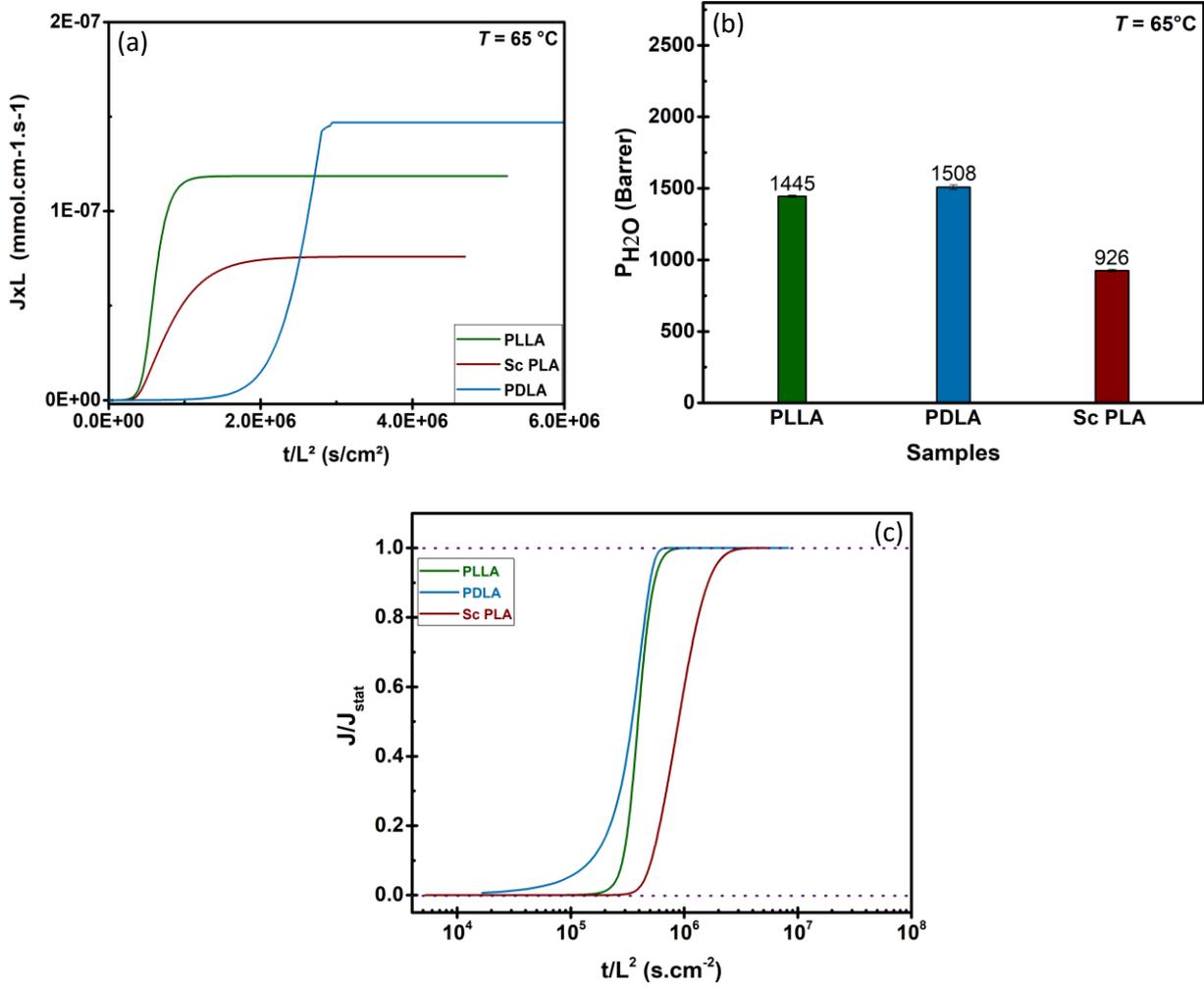
**Figure 5. 7.** (a) Evolution of the reduced flux of water permeation ( $J^*L$ ) as a function of reduced time ( $t^*L^2$ ), (b) water permeability coefficient  $P$ , and (c) normalized water permeation curves as a function of the reduced time for semicrystalline PLLA, PDLA and sc PLA at  $T_{\text{permeation}} = 45^{\circ}\text{C}$ .

45°C Samples	P (Barrer)	$D_0 \cdot 10^8$ (cm <sup>2</sup> .s <sup>-1</sup> )	$D_M \cdot 10^8$ (cm <sup>2</sup> .s <sup>-1</sup> )	$\gamma C_{\text{eq}}$	$\gamma$ (cm <sup>3</sup> .mmol <sup>-1</sup> )	$C_{\text{eq}}$ (mmol.cm <sup>-3</sup> )
PLLA	2082 ± 6	1.13 ± 0.57	16.13 ± 1.6	4.78 ± 0.37	2.59 ± 0.32	0,79 ± 0.07
PDLA	1920 ± 9	1.40 ± 0.36	14.15 ± 4.9	3.57 ± 0.05	2.23 ± 0.67	0.42 ± 0.04
Sc PLA	806 ± 4	2.14 ± 0.09	16.3 ± 3.2	2.03 ± 0.22	8.79 ± 1.77	0.24 ± 0.08

**Table 5. 4.** Water permeametric coefficients of semicrystalline PLLA, PDLA and sc PLA at  $45 \pm 1^{\circ}\text{C}$ .

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- $T_{\text{permeation}} = 65^{\circ}\text{C}$



**Figure 5. 8.** (a) Evolution of the reduced flux of water permeation ( $J \cdot L$ ) as a function of reduced time ( $t \cdot L^2$ ), (b) water permeability coefficient  $P$ , and (c) normalized water permeation curves as a function of the reduced time for semicrystalline PLLA, PDLA and sc PLA at  $T_{\text{permeation}} = 65^{\circ}\text{C}$ .

65°C Samples	$P$ (Barrer)	$D_0 \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$D_M \cdot 10^8$ ( $\text{cm}^2 \cdot \text{s}^{-1}$ )	$\gamma C_{\text{eq}}$	$\gamma$ ( $\text{cm}^3 \cdot \text{mmol}^{-1}$ )	$C_{\text{eq}}$ ( $\text{mmol} \cdot \text{cm}^{-3}$ )
PLLA	1445 ± 8	6.79 ± 0.47	15.4 ± 2.7	3.12 ± 0.26	1.24 ± 0.14	0,45 ± 0.07
PDLA	1508 ± 16	4.05 ± 0.36	12.2 ± 2.9	1.22 ± 0.08	1.08 ± 0.34	0.35 ± 0.06
Sc PLA	926 ± 8	1.2 ± 0.03	5.1 ± 1.3	1.48 ± 0.47	5.20 ± 0.87	0.28 ± 0.03

**Table 5. 5.** Water permeametric coefficients of semicrystalline PLLA, PDLA and sc PLA at  $65 \pm 1^{\circ}\text{C}$ .

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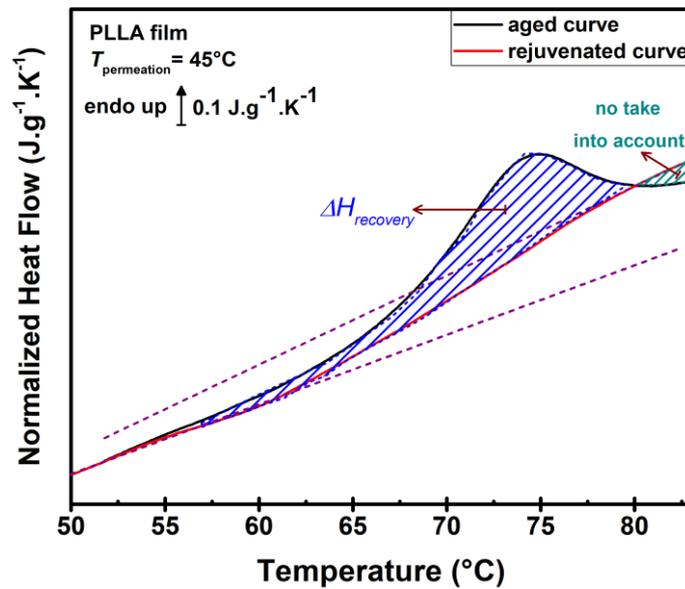
The normalized water permeation curves as a function of the reduced time are plotted to compare the kinetics of diffusion for all the samples at each  $T_{permeation}$ . An increase in the delay time for the sc PLA is observed compared to the homopolymers at all studied temperatures. In addition, one can note a decrease in the delay time observed in case of diffusion through homopolymers with  $T$  increasing, while the delay time remains practically constant in case of sc PLA (Fig 5.6 - 5.8). It is known that water can cause a plasticization effect in the polymer samples [16]. The comparison of  $D_I$  and  $D_L$  coefficients exhibits that the value of  $D_L$  is higher than the value of  $D_I$  for homopolymers (Table 5.5). Considering that the former value is related to a final period of the transient state ( $J/J_{st} = 0.62$ ), the smaller  $D_I$  value corresponds to an increase in the water diffusion during the permeation test. In general, such diffusion increase is associated with the plasticization effect of the water which leads to an increase of the material free volume [16, 18].

Temperature ( $T_{permeation}$ ) (°C)	<D> ( $\text{cm}^2 \cdot \text{s}^{-1} \cdot 10^{10}$ )			$D_{0.24}$ ( $\text{cm}^2 \cdot \text{s}^{-1} \cdot 10^{10}$ )			$D_{0.62}$ ( $\text{cm}^2 \cdot \text{s}^{-1} \cdot 10^{10}$ )		
	PLLA	PDLA	Sc PLA	PLLA	PDLA	Sc PLA	PLLA	PDLA	sc PLA
25	371	300	301	384	151	210	400	184	245
45	844	385	698	105	135	352	160	209	416
65	4712	4250	2666	1810	4110	1590	2580	4437	1820

**Table 5. 6.** The diffusion coefficient of PLLA, PDLA and sc PLA as a function of temperature.

Although according to the  $D_I$  and  $D_L$  coefficients the plasticization effect is observed. The aging phenomenon should be also taken into account in the amorphous phase organizations in case of homopolymers. Such result is totally different from the results for homopolymers found in literature. It was previously reported that thermal disturbance within the material caused an increase in the mobility of the polymer chain and, therefore, the transport phenomena were impacted by the variation of free volume resulting in increase of permeability [19, 20]. On the other hand, the aging effect could occur due to the temperature effect during the measurements. The samples were kept for purge step (drying step) under the nitrogen atmosphere at  $T_{permeation} = 45$  °C and 65 °C during more than 18 hours. Considering

the decrease in permeability coefficient, DSC analysis was performed for the studied samples in order to observe the aging effect in case of homopolymers. The DSC results (e.g. Fig 5.9) reveal that there is an endothermic peak superimposed with the glass transition which corresponds to the structural relaxation caused by aging (further results are shown in Appendix Chap 5 (A. 5.1 and 5.2)). The physical aging causes the structural relaxation towards equilibrium and results in a reduction in free volume causing lower permeability [21-25].



**Figure 5. 9.** DSC curves of studied semicrystalline PLLA after the permeation test at  $45 \pm 1^{\circ}C$ .

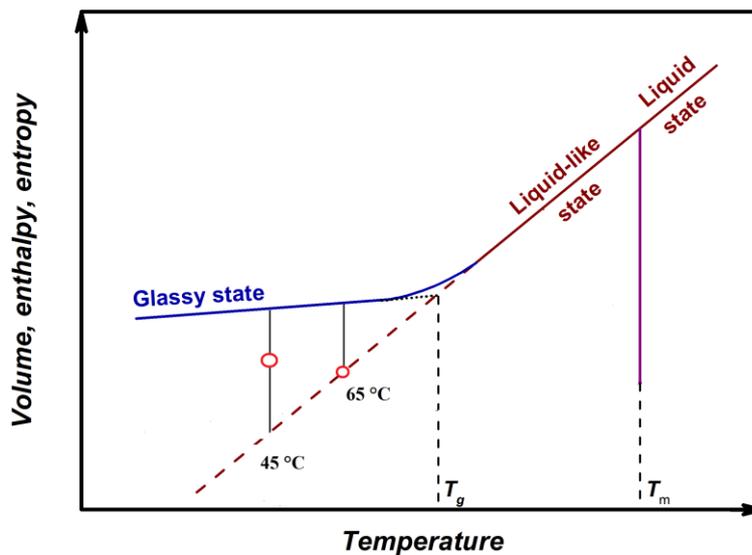
The enthalpy recovery of the structural relaxation of studied PLLA and PDLA samples were approximately calculated (shown in Fig 5.9) and results are presented in Table 5.7. The results for PLLA and PDLA were found to be 1.7 and 1.8 J/g (g of polymer) for measurement performed at 45 °C and 2.0 and 2.9 J/g (g of polymer) for measurement performed at 65 °C, respectively. Very small endothermic peak superimposed with the glass transition is observed in case of sc PLA whereas the value of the enthalpy recovery is negligible (Table 5.7) (curves are shown in Appendix Chap 5 (A. 5.1 and 5.2)). The enthalpy recovery of studied samples is normalized by the MAP content to compare with the infinite enthalpy recovery ( $\Delta H_{\infty}$ ) of wholly amorphous samples (Table 5.7). From this approximation, the enthalpy recovery values ( $\Delta H_{recovery}$  and  $\Delta H_{\infty}$ ) were compared, and, it was revealed that the MAP probably relaxed completely during the measurement performed at 65°C whereas the complete relaxation is

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not observed during the measurement performed at 45 °C. In case of measurement at 45 and 65 °C (the closest  $T$  to  $T_g$ ), the aging effect eliminates the excess free volume and increases the barrier properties. As seen from Fig 5.10 (red circles) and well known in literature, the closer to  $T_g$ , the system is quicker to reach equilibrium. So, the strong effect of aging, which is confirmed by DSC analysis, is revealed by a decrease in the permeation coefficient values during the measurements at 25 °C and up to 65 °C (Table 5.3 – 5.5).

Samples	25 °C		45 °C		65 °C		$\Delta H_\infty$
	$\Delta H_{recovery}$ (J/(g polymer))	$\Delta H_{recovery}$ (J/(g MAP))	$\Delta H_{recovery}$ (J/(g polymer))	$\Delta H_{recovery}$ (J/(g MAP))	$\Delta H_{recovery}$ (J/(g polymer))	$\Delta H_{recovery}$ (J/(g MAP))	
PLLA	ND	1.7	5	14	2	6	5
PDLA	ND	1.8	4	14	2.9	7	5
Sc PLA	ND	0.2	1	13	0.08	0.4	2

**Table 5. 7.** The enthalpy recovery values of semicrystalline PLLA, PDLA and sc PLA as a function of temperature after permeation measurements. ND: non-detected.



**Figure 5. 10.** Schematic illustration of the temperature dependence of the thermodynamic properties of the systems. Red circles present the studied samples towards equilibrium.

### 5.3.2 Gas barrier properties

The packaging materials should have high performances to be impermeable at dry state to protect the foods from the oxidation and moisture. On the other hand, fresh fruits and vegetables are the sources of biochemical reactions which consume oxygen (O<sub>2</sub>) and produce carbon dioxide (CO<sub>2</sub>) and water. Therefore, the barrier properties and selectivity of packaging materials are important for industrial applications. In this thesis, the effect of stereocomplexation on the nitrogen (N<sub>2</sub>), O<sub>2</sub> and CO<sub>2</sub> barrier properties of semicrystalline PLLA, PDLA and sc PLA films was investigated using the permeation apparatus shown in Chap. 2. The permeation and selectivity values for films were determined. Before measurement, the permeation cell was completely evacuated by applying a vacuum on both sides of the film. Then, the upstream side was provided with the gas with a pressure  $p_1$  (3 bar). The increase of pressure  $p_2$  in the calibrated downstream volume was measured using a sensitive pressure gauge linked to a data acquisition system.

The permeability can be described using Fick's first law, resulting in the permeability coefficient as follows:

$$P_i = D_i S_i \quad (5.8)$$

where  $D_i$  is the average effective diffusion coefficient (cm<sup>2</sup>/s) and  $S_i$  is the solubility coefficient (cm<sup>3</sup> (STP)/(cm<sup>3</sup> cmHg)) of component  $i$ . In addition, from the flux transient, the diffusion coefficient can be derived as follows:

$$D = \frac{L^2}{6 \times t_l} \quad (5.9)$$

where  $L$  is the sample thickness,  $D$  is the diffusion coefficient at the response time  $t_l$  called "time-lag", which corresponds to intercept of the asymptotic straight line of stationary flux with time axis.

When the permeation flux and the diffusion coefficient are determined, the permeability  $P$  and the solubility  $S$  coefficients can be calculated using the variable pressure method by assuming  $p_1 \gg p_2$  as follows [26, 27]:

$$P = \frac{J_{stat} L}{\Delta P} \quad (5.10)$$

where  $J_{stat}$  is the stationary flux, and  $\Delta P$  is the pressure difference between the upstream and downstream chambers of the permeation cell, and:

$$S = \frac{P}{D} \quad (5.11)$$

The ideal separation factor (selectivity coefficient  $\alpha (A, B)$ ) of a polymer film for gas A over gas B is given by the ratio of the gas permeability coefficients determined for gases A and B, respectively. Consequently this selectivity is the product of diffusivity and solubility selectivity:

$$\alpha (A, B) = \frac{P_A}{P_B} = \frac{D_A}{D_B} \frac{S_A}{S_B} \quad (5.12)$$

where  $D_A/D_B$  is the diffusivity selectivity and  $S_A/S_B$  is the solubility selectivity. Generally, gas diffusivity is enhanced by decreasing penetrant size, increasing polymer chain flexibility, increasing polymer fractional free volume and decreasing polymer-penetrant interactions. On the other hand, penetrant solubility is increased by increasing the condensability of the penetrant (which increases with increasing critical temperature of the gas) and increasing polymer-penetrant interactions. Therefore, the nature of the penetrant has a key role to investigate the barrier properties of studied film. Indeed, the polarizability, solubility, kinetic diameter, Van der Waals volume and critical temperature of the penetrant govern the transport properties. The characteristic parameters of used gases are presented in Table 5.8. In this thesis, the effect of the stereocomplexation on both the permeation and selectivity values for PLA-based films was investigated. The change in permeability and selectivity as a function of crystalline structure variations, free volume effects, glass transition temperature and MAP and RAF content is discussed.

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Gas	Kinetic diameter (Å)	Polarizability (cm <sup>3</sup> × 10 <sup>25</sup> )	Van der Waals volume (cm <sup>3</sup> .mol <sup>-1</sup> )	Critical temperature (°C)
N <sub>2</sub>	3.64	17.6	39.1	-147
O <sub>2</sub>	3.46	15.9	31.8	-118
CO <sub>2</sub>	3.30	26.5	42.7	31

**Table 5. 8.** Physical properties of gas [28, 29].

Due to the high permeability of homopolymers, a mechanical support, i.e. silicone film (s) is placed above the sample in order to delay the gas diffusion. The silicone film used in this study is a commercial silicone referenced as pure silicone with thickness  $L_s$  of  $0.100 \pm 0.005$  cm (purchased from Solutions Elastomères Cie, St Etienne, France). This silicone is very permeable [30]. In this case, the permeability coefficient of studied sample was calculated according to:

$$\frac{L}{P} = \sum \frac{L_i}{P_i} \quad (5.13)$$

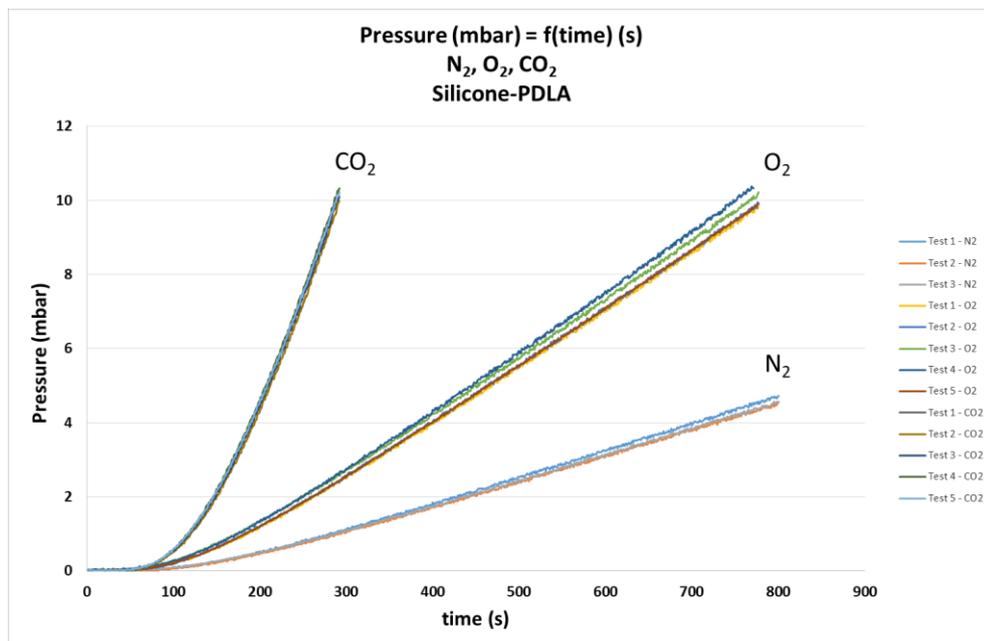
where  $L$  and  $P$  are the total thickness and the total permeability coefficient of the multilayer sample,  $L_i$  and  $P_i$  are the thickness and the permeability coefficient of each layer of the multilayer system, respectively. The quantity  $R = L/P$  is the resistance of the sample to gas permeation. In the case of a multilayer sample, the total resistance is the sum of the resistance ( $R_i = L_i/P_i$ ) of each layer [31]. The resistance  $R_s = L_s/P_s$  of the silicone sample and the resistance of the multilayer sample ( $R_{su} = L_{su}/P_{su}$  silicone/unknown sample) can be determined. The resistance  $L_u/P_u$  of the unknown film can be deduced from the global resistance by:

$$L_{SU}/P_{SU} = L_S/P_S + L_U/P_U \quad (5.14)$$

$$P_U = L_U/(L_{SU}/P_{SU} - L_S/P_S) \quad (5.15)$$

The gas permeation measurements were carried out firstly for pure silicone films. The results are presented in Appendix Chap 5 (A. 5.3, A. 5.4 and A. 5.5). The increase of the pressure  $p_2$  as a function of time is presented for all tested gases (N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>). Considering the gas permeability dependence on the nature and size of the penetrant, and also on the nature of the polymer, D.W. Van Krevelen [32] proposed the permeability coefficients of N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>

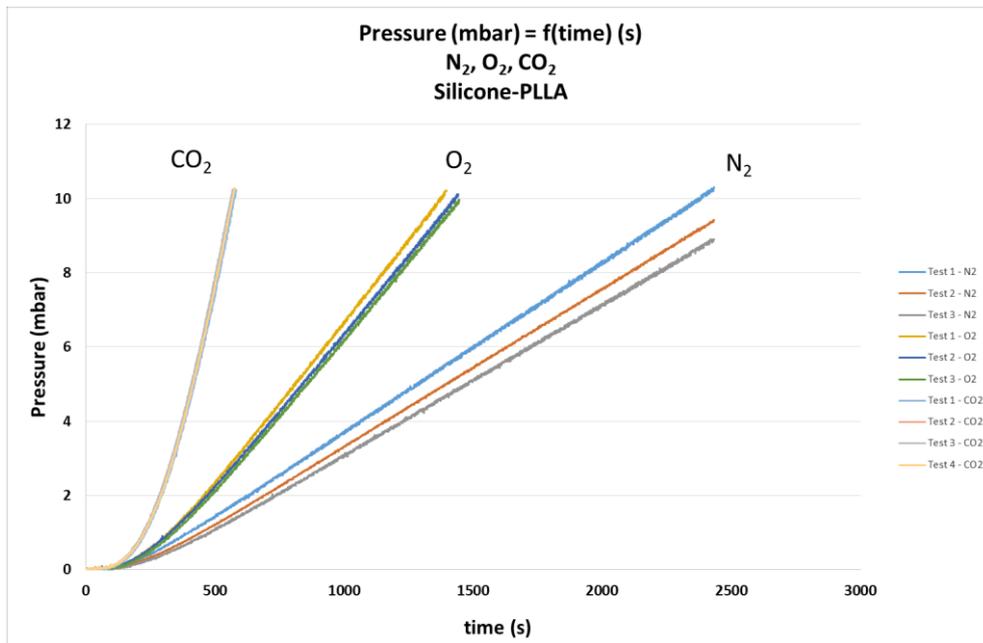
gases follow the order according to their kinetic diameter as:  $P(N_2) < P(O_2) < P(CO_2)$ . The same order was observed in case of the silicone film. Permeability coefficients and the values of selectivity ( $\alpha$ ) of the silicone film towards  $N_2$ ,  $O_2$  and  $CO_2$  were calculated and results are presented in Table A. 5.4 and Table A. 5.5. After silicone film, the silicone-PDLA bilayer system was studied to calculate the permeability coefficient of PDLA film. The pressure curves (Fig 5.11) reveal that the permeability coefficients follow the same order as proposed depending on the kinetic diameters of penetrants ( $P(N_2) < P(O_2) < P(CO_2)$ ). Furthermore, the experimental pressure curves for each gas are practically all superimposed (Fig 5.11) that indicates an excellent reproducibility of the measurements.



**Figure 5. 11.** Pressure as a function of time for silicone-PDLA system.

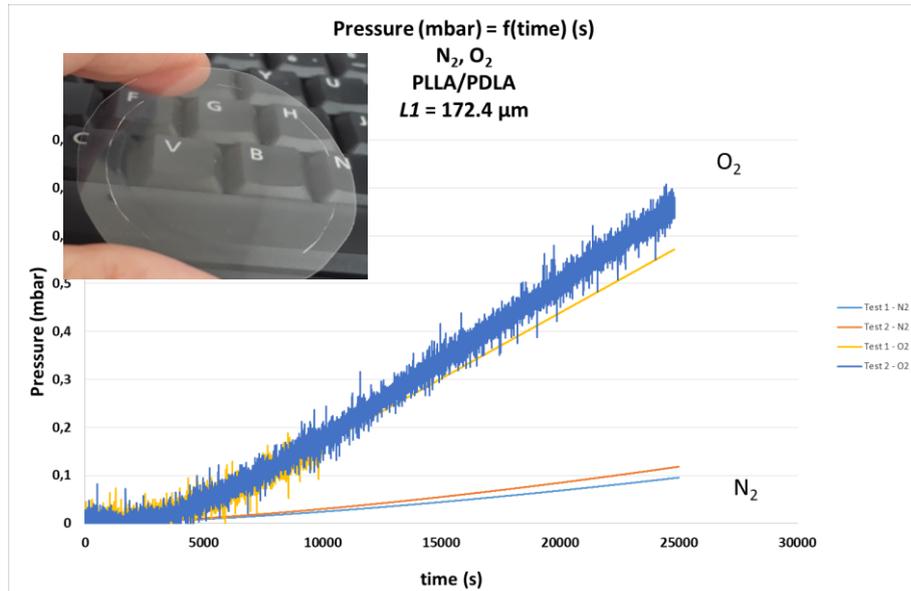
The permeation coefficient and the selectivity of PDLA film can be calculated by applying the law of series of resistance to diffusion (Eq 5.15) (Table 5.9). The same technique was applied also for PLLA film to calculate the permeability parameters towards  $N_2$ ,  $O_2$  and  $CO_2$ . The experimental curves for each gas are practically all superimposable (Fig 5.12), nevertheless we can note that the results with the nitrogen gas are more dispersed. Such behavior can be explained by the densification of the polymer film caused by specific interactions between the polymer film and  $N_2$ . Indeed, permeation of larger penetrants

molecules (i.e.  $N_2$ ) is more affected by changes in free volume than permeation of small molecule (i.e.  $CO_2$ ) [33]. The permeation parameters of PLLA film can be also calculated by the law of series of resistance to diffusion (Eq 5.15) (Table 5.9).

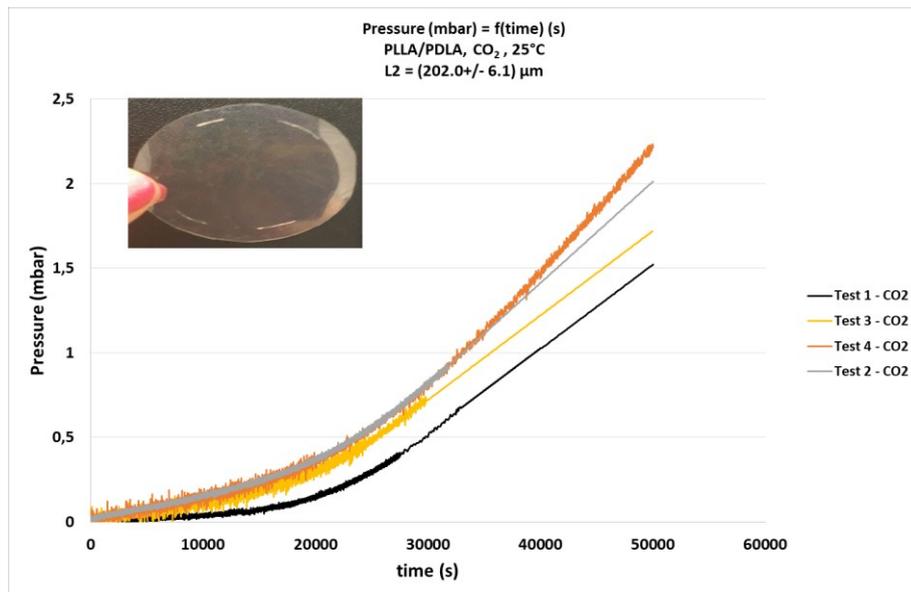


**Figure 5. 12.** Pressure as a function of time for silicone-PLLA system.

The use of silicone film was not necessary for sc PLA. However, the high mechanical fragility of sc PLA was a problem because of the film crack caused by the vacuum. The experimental curves (Fig 5.13) reveal that the results are not perfectly superimposable. It was impossible to study  $CO_2$  permeation with the same sc PLA sample, because of the sample cracking during the drying step. The permeation tests with  $CO_2$  have been realized with another sc PLA sample, and we supposed that the sample was cracked between the 3<sup>rd</sup> and 4<sup>th</sup> test (Fig 5.14). And, really the photo of sc PLA samples indicates that the samples are damaged during the measurements. Therefore, only the results of first tests were taken into account.



**Figure 5. 13.** Pressure as a function of time for sc PLA film towards N<sub>2</sub> and O<sub>2</sub>.



**Figure 5. 14.** Pressure as a function of time for sc PLA film towards CO<sub>2</sub>.

The permeability coefficient of all studied samples is presented in Table 5.9. As can be seen from the results, whatever the studied film, the same tendency is observed for the order of permeability coefficient ( $P(\text{CO}_2) > P(\text{O}_2) > P(\text{N}_2)$ ), i.e. according to the gas kinetic diameter. Also whatever the penetrant gas, the sc PLA has lower permeability coefficient than that for homopolymers. Indeed, this result can be explained by the fact that the permeability

coefficient is a result of the diffusion coefficient (which corresponds to the mobility of the gas through the film) and the solubility coefficient (which reflects the capacity of a molecule to condense in the polymer). So, due to the lower kinetic diameter (3.30 Å) and highest critical temperature (31 °C), CO<sub>2</sub> has the highest permeability coefficient value among all gases studied. The presence of polar ester oxygens in the polymer chain favors the solubility of quadrupolar CO<sub>2</sub> over the other non-polar gases [34-36].

As reported in literature, the presence of crystals is a predominant factor to cause more tortuosity in the film and to make it less permeable. In our case, this factor was not taken into consideration due to the similar crystallinity degree of studied films (Table 5.1). Nassar et al. [37] reported that the RAF content in the semicrystalline samples has strong influence on the barrier properties. This is due to RAF insufficient decoupling from the crystalline phase that provides an accelerated pathway for diffusion and, thus increases permeability. Therefore, we focused our attention on the influence of the crystals type ( $\alpha$  and sc) and the RAF content in the semicrystalline films. In our case, the higher RAF content is observed in case of sc PLA (Table 5.1).

Permeability coefficient <i>P</i> (Barrer)	N <sub>2</sub>	O <sub>2</sub>	CO <sub>2</sub>
<b>PDLA</b>	138 ± 21	339 ± 44	2081 ± 187
<b>PLLA</b>	38 ± 12	71 ± 25	283 ± 22
<b>Sc PLA</b>	0.024 ± 0.004	0.14 ± 0.02	0.28 ± 0.04

**Table 5. 9.** The permeability coefficient of studied films towards N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>.

However, sc PLA is found to be more barrier than homopolymers films. Indeed, the measurement temperature ( $T = 25 \pm 1$  °C) is below the glass transition temperature of studied samples (Table 5.1). Therefore, film samples are in the glassy state, so the polymer chains are rigid and less mobile. In addition, sc PLA has the lowest glass transition temperature (about > 5 °C), nevertheless sc PLA presents better barrier properties whatever the penetrant gases. To better understand the transport properties of the studied samples, the variations of the different permeametric coefficients (i.e. diffusion and solubility coefficients) were also investigated. The diffusion and solubility coefficient values are presented in Table 5.10 and

5.11, respectively. The diffusivity of a gas is related to the mobility of this gas within the polymer film. The value of the diffusion coefficient will therefore mainly depend on the size of the penetrating molecules. The smaller the volume of the gas molecule, the easier it will diffuse through the film (i.e. a molecule with a larger volume will be slowed down and, therefore, will have more difficulty to move through the polymer chains). So, according to the Van der Waals volumes of gases, i.e.  $V_{O_2} = 31.8 \text{ cm}^3 \cdot \text{mol}^{-1} < V_{N_2} = 39.1 \text{ cm}^3 \cdot \text{mol}^{-1} < V_{CO_2} = 42.7 \text{ cm}^3 \cdot \text{mol}^{-1}$  (Table 5.8) the values of the diffusion coefficient should have such tendency as  $D(CO_2) > D(N_2) > D(O_2)$ . As can be seen from Table 5. 10, only PDLA film has this tendency, whereas PLLA has tendency as  $D(N_2) > D(O_2) > D(CO_2)$  which can be explained by the condensation effect of  $O_2$  resulting in the decrease in diffusion. In addition, sc PLA presents inverse tendency with the Van der Waals volume as  $D(CO_2) < D(N_2) < D(O_2)$ . Due to the steric hindrance in sc PLA, the results show that the bigger the size of the penetrant molecule, the longer time will be required to cross the film, and the diffusion will be slowed down. The diffusion coefficients evolution exhibit that the transport of gases through these films is not done in the same way. This difference can be related to the different crystalline structure (sc and  $\alpha$ ) between sc PLA and homopolymers and, therefore, related to the free volume and RAF content.

Diffusion coefficient . $10^8$ $D \text{ (cm}^2 \cdot \text{s}^{-1}\text{)}$	$N_2$	$O_2$	$CO_2$
<b>PDLA</b>	$1169 \pm 32$	$1397 \pm 55$	$1424 \pm 27$
<b>PLLA</b>	$1025 \pm 120$	$877 \pm 34$	$796 \pm 19$
<b>Sc PLA</b>	$0.57 \pm 0.05$	$1.15 \pm 0.13$	$0.34 \pm 0.12$

**Table 5. 10.** The diffusion coefficient of studied films towards  $N_2$ ,  $O_2$  and  $CO_2$ .

The different diffusion coefficients values were obtained for the studied samples, therefore, the lower permeability coefficient value in the case of sc PLA may be correlated with the solubility coefficients values. The obtained solubility coefficients of studied samples are presented in Table 5. 11. The solubility coefficient of polymers has an evolution according to the critical temperatures of the penetrant molecules as follows:  $S(N_2) < S(O_2) < S(CO_2)$ , ( $T_c(N_2) < T_c(O_2) < T_c(CO_2)$ ) (Table 5.8) [38, 39]. As can be seen from Table 5.11, the solubility

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coefficient values of studied films present good accordance with literature. In the case of sc PLA, the lowest solubility values are observed suggesting an improvement of the barrier properties, except for CO<sub>2</sub>. The higher CO<sub>2</sub> solubility coefficient can be explained by the RAF content which effects the solubility coefficient by increasing the free volume.

Solubility coefficient . 10 <sup>2</sup> S (cm <sup>3</sup> (STP).cm <sup>-3</sup> cmHg)	N <sub>2</sub>	O <sub>2</sub>	CO <sub>2</sub>
PDLA	0.18 ± 0.01	0.32 ± 0.01	1.32 ± 0.04
PLLA	0.13 ± 0.01	0.29 ± 0.01	1.29 ± 0.06
Sc PLA	0.05 ± 0.01	0.12 ± 0.03	1.36 ± 0.31

**Table 5. 11.** The solubility coefficient of studied films towards N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>.

The efficiency of the studied films to separate the components in the penetrants mixture can be defined by the selectivity parameters presented in Table 5.12. Although the strong decrease in permeability is observed in the case of for sc PLA for both water and gases, the significant selectivity is observed only for systems containing water. Therefore, one can note that stereocomplexation causes more important decrease in gas barrier properties of film compare to the water barrier properties which must be taking into account for food packaging. In addition, due to the biochemical reaction products, decreasing the respiratory intensity is always aimed for the longer life-time, which can be obtained by the reduction of partial pressure of oxygen in the packaging materials. Moreover, the presence of H<sub>2</sub>O and CO<sub>2</sub> leads to a bad taste and smell in the packed products. Therefore, high H<sub>2</sub>O / O<sub>2</sub> permselectivity and the low O<sub>2</sub> permeability is an optimum condition to reduce the respiratory intensity and the anaerobic fermentation rate to have longer life-time. The result of permselectivity (Table 5.12) shows that stereocomplexation not only improves barrier properties, but also achieves the  $\alpha_{H_2O / O_2}$  value of PLA-based materials for packaging applications. The trade-off between the H<sub>2</sub>O and O<sub>2</sub> permeability and the ideal separation factor  $\alpha_{H_2O / O_2}$  was found in case of sc PLA.

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Selectivity ( $\alpha_{A/B}$ )	CO <sub>2</sub> / O <sub>2</sub>	CO <sub>2</sub> / N <sub>2</sub>	O <sub>2</sub> /N <sub>2</sub>	H <sub>2</sub> O / O <sub>2</sub>	H <sub>2</sub> O / N <sub>2</sub>	H <sub>2</sub> O / CO <sub>2</sub>
<b>PDLA</b>	6	15	2	6	15	1
<b>PLLA</b>	4	7	2	34	65	9
<b>Sc PLA</b>	2	12	6	5630	31667	2713

**Table 5. 12.** The selectivity parameters of studied films.

Moreover, the sc PLA results show better improvements and/or are in good accordance with previous works to achieve better barrier properties of PLA-based polymers. For example, Picard et al [40] reported that the barrier properties of PLA nanocomposites film can be improved by increasing the content of organo-modified montmorillonite (O-MMT). In that work, crystallinity degree was increased up to 35% by annealing time and temperature. For the maximum crystallinity degree value of annealed and charged (4 wt %) PLA samples ( $X_c = 46\%$ ), the O<sub>2</sub> permeability coefficient of PLA composite was calculated as  $0.23 \pm 0.03$  Barrer. In another work, Pinto et al. [41] studied nanocomposite thin films of PLA which were produced by incorporating small amounts (0.2 to 1 wt%) of graphene oxide (GO) and graphene nanoplatelets (GNP). The permeabilities towards O<sub>2</sub> are decreased from 3.76 Barrer for PLA to 1.49 and 1.30 Barrer; and the N<sub>2</sub> permeability coefficients are decreased from 1.10 Barrer for PLA to 0.502 and 0.237 Barrer for nanocomposite film with increasing the percentage GO and GNP up to 0.6 wt%, respectively. Furthermore, Lehermeier et al. [42] investigated barrier properties of amorphous PLA and biaxially oriented semicrystalline PLA film. They proposed that polymer chain branching and small changes in L:D stereochemical content of PLA have no influence on the permeation properties and the most promoted influence is caused by the crystallinity degree. The sample prepared in that work [42] has the crystallinity degree  $X_c = 18\%$  and the permeability coefficients of N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub> were determined to be 1.3, 3.3 and 10.2 Barrer, respectively. So, one can say that the results of the sc PLA studied in this thesis show superior barrier properties of PLA-based material compare to the previous works. Furthermore, sc PLA possesses enhanced barrier properties compared to the petroleum-based polymers usually used in the food packaging (Table 5.13).

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Samples	N <sub>2</sub>	O <sub>2</sub>	CO <sub>2</sub>
Sc PLA	0.03	0.14	0.28
LDPE	1.9	6.9	28.0
PS	2.2	2.6	10.5
PET	0.008	0.04	0.2

**Table 5. 13.** The permeability coefficient of sc PLA compared to the petroleum-based polymers (LDPE: low-density polyethylene, PS: polystyrene, PET: polyethylene terephthalate) [42].

### Conclusion

Nowadays, a market of packaging requires more eco-friendly materials with high performances, i.e. strong mechanical and barrier properties. In this thesis, we proposed facile method in order to improve the mechanical and transport properties of PLA, namely the stereocomplex formation between PLLA and PDLA. The water resistance of amorphous and semicrystalline PLA-based film and the permeation of nitrogen, oxygen and carbon dioxide in semicrystalline samples were studied to examine the influence of the stereocomplexation on transport properties. The presence of pure homo- or sc crystals was confirmed by XRD and DSC analysis. The morphology of the obtained films was investigated by SEM analysis. SEM results show that the homopolymers films exhibit granular structure, while sc PLA films reveal a smooth compact structure. Furthermore, no porosity is observed for all the film samples.

Superior mechanical properties have been achieved by stereocomplexation of homopolymers through formation of stereocomplex crystals which are formed by intermolecular crosslinking. The barrier properties of poly(lactide) were also enhanced by stereocomplexation.

One should note that although there is no difference on the glass transition properties of amorphous samples (i.e. the same glass transition, molecular mobility and CRR size), the water barrier properties of amorphous PLA-based sample can be improved by stereocomplexation. Furthermore, the value of water permeability coefficient of amorphous sc PLA at 25°C is close to the value of water permeability coefficient of semicrystalline ( $X_c = 48\%$ ) homopolymers.

The transport properties of semicrystalline pure PLLA, PDLA and sc PLA were investigated by water permeation measurements as a function of the temperature. The water permeation results reveal an important barrier improvement for sc PLA - up to 70% as compared to the homopolymers. It is found that even if the molecular structure of samples is the same, the stereocomplexation creates the macromolecular reorganization in PLA-based films. Moreover, one can note that the type of crystals is the predominant factor on the barrier properties compared to the RAF content (i.e. although sc PLA has higher RAF content than homopolymers, it is still more barrier than homopolymers). It is also very interesting that the barrier properties of homopolymers are found to increase with the  $T$  increase due to the aging effect revealed in case of homopolymers.

The gas permeation results clearly show that stereocomplexation not only enhances the water barrier properties, but also improves the gas barrier properties. The results of sc PLA reveal that the proposed new method can be applied to the biobased homopolymers, which have different tacticity, to increase the competition with non-ecofriendly petroleum-based polymers used in food packaging applications. From that point, the investigation of the relation between the microstructure of PLA and the water and gas barrier performance is still a key in the improvement of the transport properties of PLA towards different permeants. Nevertheless, the improved barrier properties of sc PLA (PLLA/PDLA) films through enhanced stereocomplexation may comply with the requirement for high temperature engineering and packaging application.

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### CONCLUSIONS & PROSPECTS

In recent years, PLA-based materials have attracted great interest as a biobased ecofriendly polyester to replace petroleum-based plastic used in many applications (i.e. biomedical devices, tissue engineering, and in packaging). Therefore, the new production strategy, called stereocomplexation, is developed to obtain a material with better physical properties. Moreover, the behavior of the amorphous phase and the transport properties of stereocomplex PLA was still enigmatic topic. In that concern, the goal of this thesis is to have better understanding and advanced investigation of the amorphous phase properties and the evolution of the barrier properties of stereocomplex PLA with the aim of improving the knowledge about sc PLA and stereocomplexation reactions.

The first part of this thesis was devoted to optimization of the elaboration method of stereocomplex PLA with pure stereocomplex crystals as a crystalline structure. For this purpose, stereocomplex PLA was elaborated by extrusion process (at different extrusion temperature and extrusion time) and by solution casting (with different polymer concentrations and homopolymers' ratio). The thermal and microstructural analyses of the obtained films reveal that:

- ♦ extrusion process allows decreasing the thermal degradation temperature of stereocomplex PLA with an increasing extrusion temperature and time, whereas the solution casting method has no influence on the thermal behavior of sc PLA;
- ♦ extrusion temperature ( $T_{ext}$ ) is found to have influence on the stereocomplex formation as pure and stable stereocomplex crystals were obtained at  $T_{ext} = 220^{\circ}\text{C}$ ;
- ♦ the polymer concentration has slight influence on the stereocomplexation and the homopolymer presence. Besides, the obtained stereocomplex crystals by solution casting are not stable;
- ♦ the ratio of homopolymers has significant influence on the stereocomplex formation and the found optimum conditions of blending ratio are 60:40 and 50:50 for PLLA:PDLA;

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- ♦ semicrystalline sc PLA obtained by extrusion process presents 2-phase model ( $X_{MAP} + X_C = 1$ ), whereas semicrystalline sc PLA obtained by solution casting reveals 3-phase model ( $X_{MAP} + X_C + X_{RAF} = 1$ ) due to the different crystallization temperature which is higher in the case of extrusion process.

In the second part of this study, the objective was to investigate the chirality/tacticity impact on the molecular dynamics of amorphous stereocomplex PLA. The physical aging study was carried out by FSC in order to examine the glass transition parameters and the structural relaxation of amorphous stereocomplex PLA. The dynamic parameters associated to the  $\alpha$  relaxation process, i.e. the relaxation time, the dielectric strength, and the fragility, have been investigated by DRS. The dynamic heterogeneities and the cooperative length scales at the dynamic glass transition have been studied by combining DRS and MT-DSC techniques. According to the molecular dynamic investigation, one may conclude that:

- ♦ amorphous stereocomplex PLA has the same glass transition temperature as amorphous homopolymers;
- ♦ the enthalpy recovery has a single step decay towards equilibrium and the relaxation kinetics of the glass (PLLA, PDLA and sc PLA) is independent on the tacticity in stereocomplex structure;
- ♦ the dielectric properties are independent on the tacticity presence;
- ♦ the cooperativity length  $\xi_\alpha$  and cooperativity number  $N_\alpha$  have no dependence on the chemical structure (i.e. chirality) in the case of amorphous samples;
- ♦ through the  $\beta$ -process analysis it is found that the local movements of polymer chain are independent from the stereocomplexation in case of amorphous samples. However, different crystalline phase is obtained during stereocomplexation.

Our aim was also to investigate the influence of the stereocomplexation on the transport properties of PLA. Therefore, last part of this thesis is focused on the analysis of the barrier properties towards water and gases ( $N_2$ ,  $O_2$ , and  $CO_2$ ). Special attention was paid to the water permeation of semicrystalline stereocomplex PLA as a function of temperature.

According to the permeation measurements results, the improvements of barrier properties of PLA can be presented as follows:

- ♦ the glass transition properties and molecular dynamics of amorphous samples was found to be same for homopolymers and sc PLA and the stereocomplex presence improves the barrier properties of amorphous stereocomplex PLA;
- ♦ in case of semicrystalline samples, an important barrier improvement for stereocomplex PLA was observed; i.e. up to 70% as compared to the homopolymers;
- ♦ stereocomplex PLA shows better barrier properties with increasing the temperature of permeation measurements;
- ♦ the aging effect on the barrier properties of homopolymers was observed as decrease in the values of permeation coefficient with increasing measurement temperature;
- ♦ the gas barrier properties are also enhanced by stereocomplexation, thus stereocomplexation can be used as a new method to polymers synthesis for food packaging applications.

Future work can be directed towards the production optimization of stereocomplex PLA films through the extrusion process if it were to be targeted for packaging applications. From this optimization, different amorphous fractions could be obtained and analyzed (i.e. dielectric properties, barrier properties, etc.) for its barrier properties as the function of temperature. A complete gas permeation measurement as the function of temperature for both amorphous and semicrystalline samples can also be carried out to study stereocomplexation effect on the transport properties of PLA. Rheology analysis at high temperature can be also performed to investigate mechanical properties of stereocomplex PLA.

The study of molecular dynamics as a function of frequency and temperature for semicrystalline samples is also crucial to understand the chirality effect of stereocomplex PLA. High pressure DRS analysis could be utilized to reveal the piezoelectric properties of stereocomplex PLA. Over the last decade, there is an increase interest towards the piezo-

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response of biobased polymers as compared to other synthetic piezo-materials (e.g. ceramics). Besides packaging applications, the understanding of piezoelectricity is useful in applications that involve separation membranes, pressure sensors, energy harvester and transparent nanogenerator, and ultrasound-based systems (microphones, echolocation).



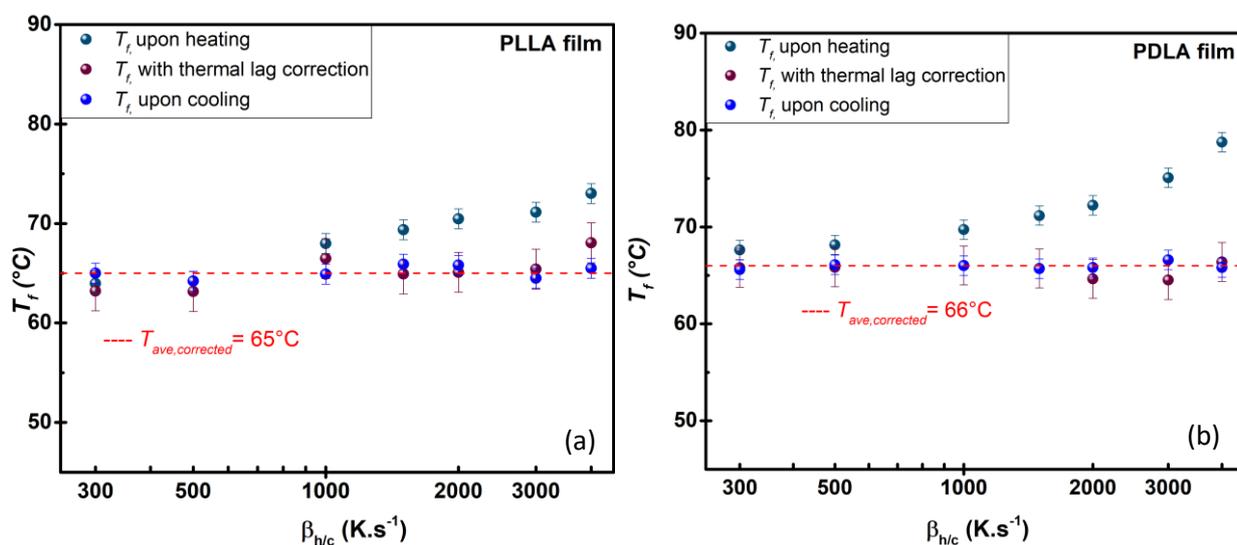


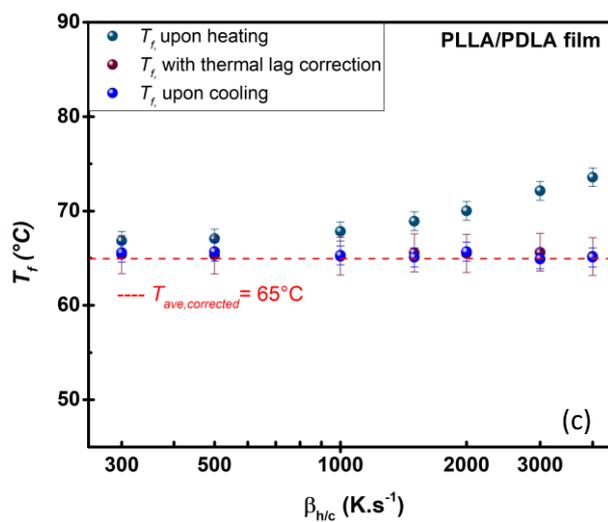
APPENDIX

Chapter 4

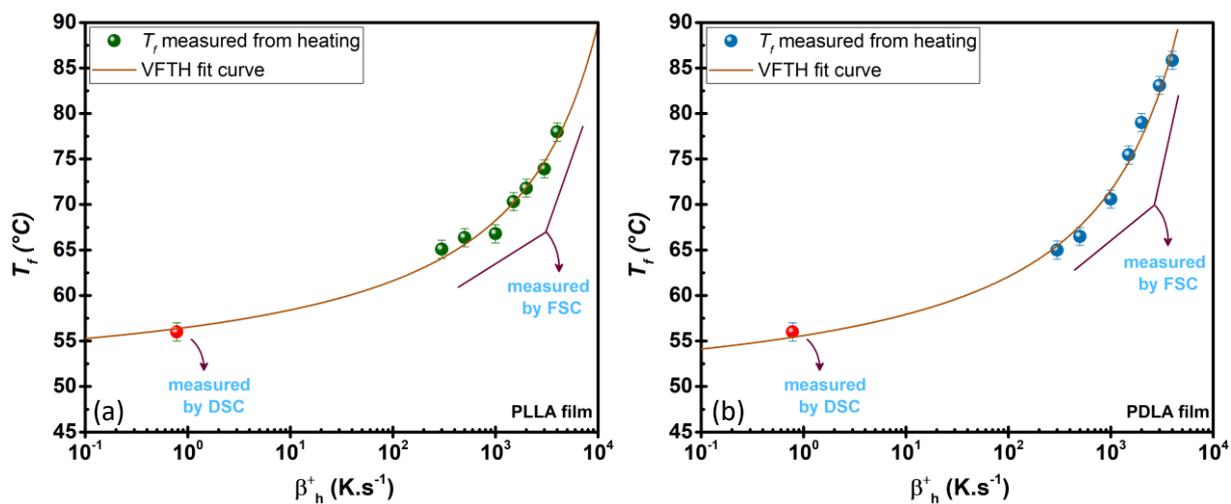
$ \beta_c  = \beta_h$ (Ks <sup>-1</sup> )	$\delta T_L$			$T_f$ from heating (°C)			$T_f$ corrected from heating (°C)			$T_f$ from cooling (°C)		
	(A)	(B)	(C)	(A)	(B)	(C)	(A)	(B)	(C)	(A)	(B)	(C)
300	0.8	1.8	1.5	64.0	67.6	66.8	63.2	65.8	65.4	65.0	65.6	65.6
500	1.0	2.3	1.8	64.2	68.1	67.1	63.2	65.8	65.3	64.2	66.1	65.7
1000	1.5	3.7	2.6	68.0	69.7	67.8	66.5	66.0	65.2	64.9	66.0	65.3
1500	4.4	5.5	3.3	69.4	71.2	68.9	64.9	65.7	65.6	65.9	65.7	65.1
2000	5.4	7.6	4.5	70.5	72.2	70.0	65.1	64.6	65.5	65.8	65.8	65.7
3000	5.7	10.6	6.5	71.1	75.1	72.1	65.4	64.5	65.6	64.5	66.6	64.9
4000	5.0	12.4	8.4	73.0	78.8	73.6	68.1	66.4	65.2	65.5	65.8	65.1

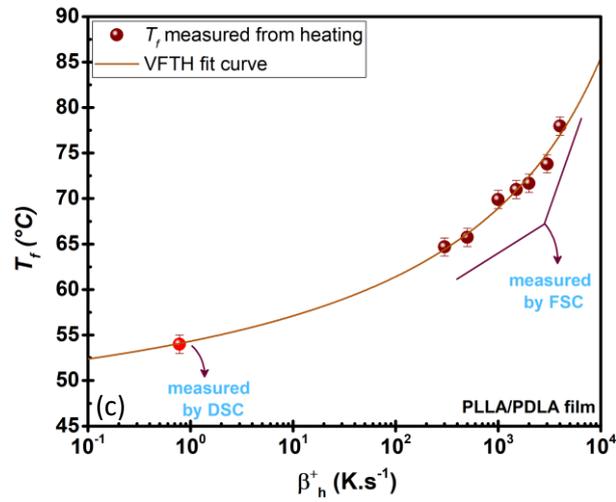
A. 4. 1. The measured and corrected values of the total thermal lag and the fictive temperatures for the different scanning rates for (A) PLLA, (B) PDLA, (C) sc PLA.





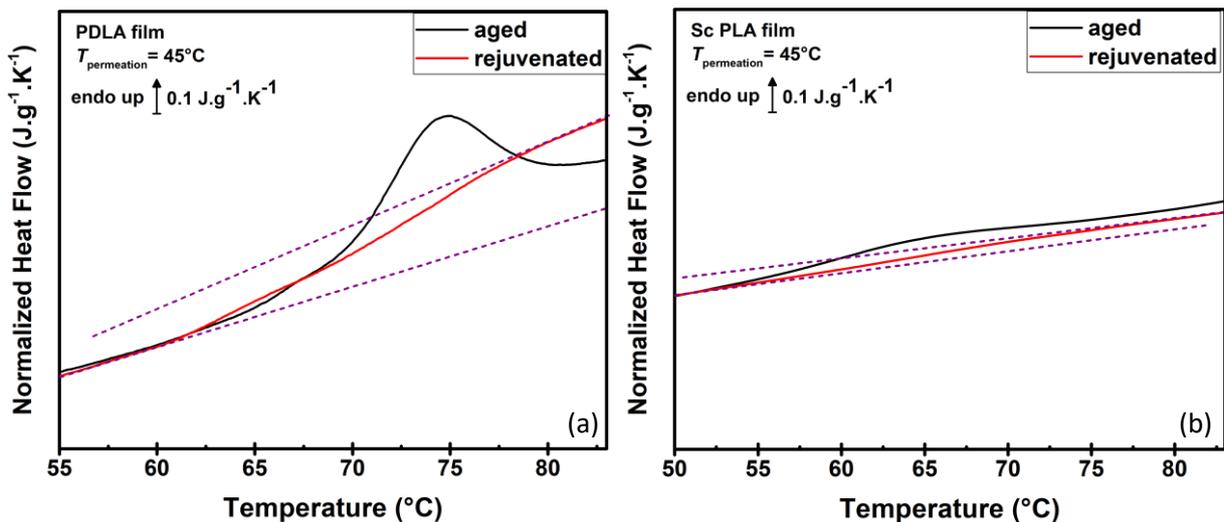
A. 4. 2. Evolution of the fictive temperature as a function of equivalent scanning rate for amorphous samples: (a) PLLA, (b) PDLA and (c) sc PLA.



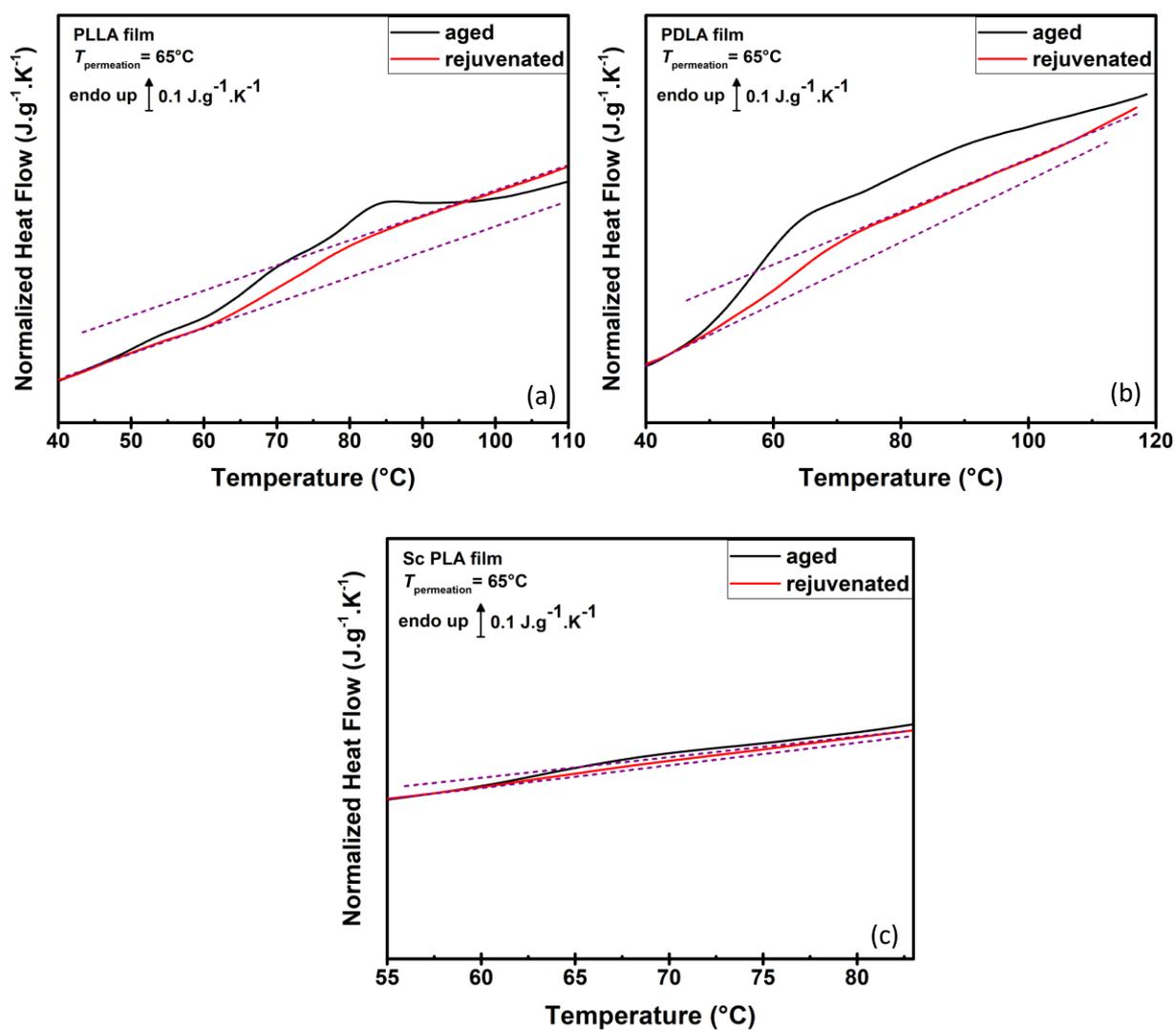


**A. 4. 3.** Evolution of the fictive temperature measured upon heating as a function of the scanning rate  $|\beta_c| = \beta_h$  for (a) PLLA, (b) PDLA and (c) PLLA/PDLA. The red-colored symbols present the fictive temperature determined from DSC at  $|\beta_c| = \beta_h = 0.78 \text{ K.s}^{-1}$ . The other results present the fictive temperature determined from FSC at  $|\beta_c| = \beta_h$  ranging from  $300 \text{ K.s}^{-1}$  up to  $4000 \text{ K.s}^{-1}$ . The solid lines correspond to the VFTH fit.

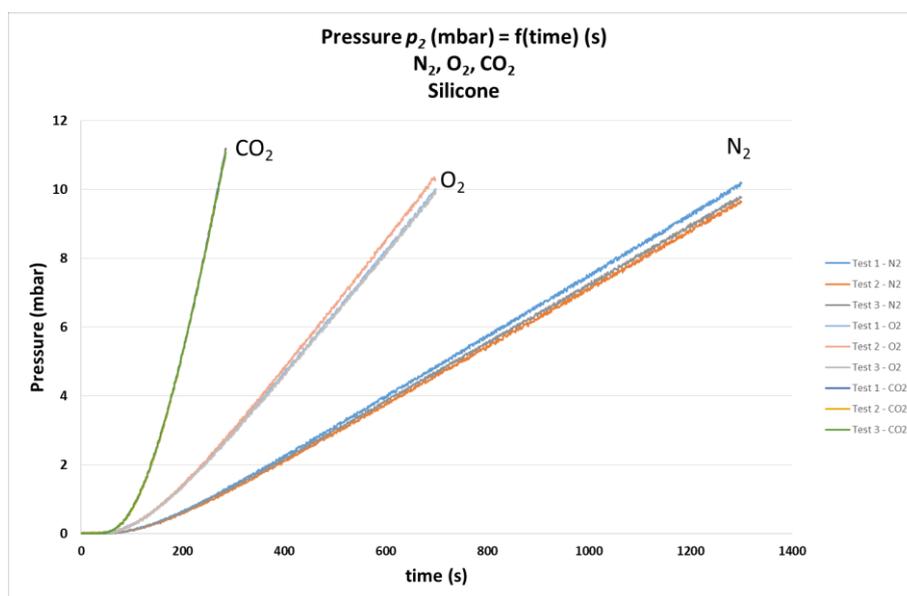
Chapter 5



**A. 5. 1.** DSC curves of studied semicrystalline: (a) PDLA and (b) sc PLA after the permeation test at  $45 \pm 1^\circ\text{C}$ .



A. 5. 2. DSC curves of studied semicrystalline: (a) PLLA, (b) PDLA and (c) sc PLA after the permeation test at  $65 \pm 1^{\circ}\text{C}$ .



A. 5. 3. Pressure as a function of time for silicone film.

Permeability coefficient P (Barrer)	N <sub>2</sub>	O <sub>2</sub>	CO <sub>2</sub>
Silicone	215 ± 1.7	460 ± 10	1853 ± 6

A. 5. 4. The permeability coefficient of silicone film towards N<sub>2</sub>, O<sub>2</sub> and CO<sub>2</sub>.

Selectivity	O <sub>2</sub> /N <sub>2</sub>	CO <sub>2</sub> / N <sub>2</sub>	CO <sub>2</sub> / O <sub>2</sub>
α	2.1	8.6	1.0

A. 5. 5. The selectivity of silicone film.







## **Analyse Thermique Avancée et Propriétés de Transport de Matériaux PolyLactide Stéréocomplexe**

L'originalité de ce travail repose sur une approche physique et physico-chimique des propriétés de mélanges de polylactides de chiralité différentes (poly L-lactique acide et poly D-lactique acide) et sur l'influence de cette chiralité sur les propriétés des phases amorphes de ces matériaux. Les matériaux sont des mélanges élaborés à partir de deux homopolymères (PLLA et PDLA) selon deux méthodes ; coulée en solution ou par extrusion. Nous avons étudié des matériaux totalement amorphes et cristallisés de façon isotrope afin de générer une phase amorphe plus ou moins confinée. Il est montré que le mélange pouvant donner naissance à une phase cristalline stéréocomplexe ne peut être obtenu que dans certaines conditions. Les résultats des analyses thermiques et de perméation ont montré que le mélange PLLA/PDLA améliore certaines propriétés du matériau, notamment des propriétés barrières à l'eau et aux gaz plus élevées par rapport aux homopolymères parents. Afin d'étudier la mobilité moléculaire des phases amorphes, du vieillissement physique et de la relaxation structural (relaxation  $\alpha$  et  $\beta$ ), le concept de Région de Réarrangement Coopératif (CRR) a été appliqué. Il a été montré que les phases amorphes des homopolymères et du mélange ont exactement les mêmes propriétés à la transition vitreuse et dans l'état vitreux lorsque les matériaux sont totalement amorphes.

**Mots-clés:** stéréocomplexation, PLA stéréocomplexe, propriétés de transport, analyse thermique, dynamique moléculaire, relaxation structurale, vieillissement physique, Spectroscopie de Relaxation Diélectrique, Calorimétrie à Balayage Rapide, Perméation

### **Advanced Thermal Analysis and Transport Properties of Stereocomplex PolyLactide**

The originality of this work is based on analysis of physical and physicochemical properties of polylactide mixtures of different chirality (poly L-lactic acid and poly D-lactic acid) and on the influence of the chirality on the amorphous phase's properties. The materials mixtures are elaborated from two homopolymers (PLLA and PDLA) according to two methods; solution casting or extrusion. Totally amorphous and isotropically crystallized materials with more or less confined amorphous phase were studied. It is shown that a stereocomplex crystalline phase can be obtained only under certain experimental conditions. The results of the thermal and permeation analyzes showed that the PLLA / PDLA mixture improved certain properties of the material, namely higher barrier properties towards liquid water and gases were obtained compared to parent homopolymers. In order to study the molecular mobility of amorphous phases, physical aging and structural relaxation ( $\alpha$  and  $\beta$  relaxation), the Cooperative Rearrangement Region (CRR) concept has been applied. It has been shown that the amorphous phases of the homopolymers and the mixture have exactly the same properties at the glass transition and in the vitreous state when the materials are totally amorphous.

**Keywords:** stereocomplexation, stereocomplex PLA, transport properties, thermal analysis, molecular dynamic, structural relaxation, physical aging, Dielectric Relaxation Spectroscopy, Fast Scanning Calorimetry, Permeation