



Contribution ID: 172

Type: **Orale**

Effect of Molecular Weights on the Physical Properties and Microstructure of Poly(lactic acid)

Wednesday, 10 July 2019 15:30 (15 minutes)

A three-phase microstructure involving crystalline, rigid and mobile amorphous regions has been successfully used to describe the macromolecular arrangements in poly(lactic acid) (PLA). The establishment of a rigid amorphous fraction, in particular, is dependent on the molecular motions of polymer chains during crystallization; Our aim is to understand the contribution of the size of the polymer chains (related to their number average molar mass, M_n) on its molecular structure during crystallization. To achieve this, a commercial grade of PLA was hydrolyzed by extrusion in controlled conditions of time, temperature and water content. The glass transition temperatures of these fractions obey the Flory-Fox model and reach a plateau beyond ~ 20000 g/mol. Preliminary measurements performed by Differential Scanning Calorimeter (DSC) showed that, as M_n decreases and reaches values lower than ~ 40000 g/mol, cold crystallization starts to occur. As M_n further decreases, the melting peaks associated to cold-crystallized domains gradually shifts towards lower temperatures. PLA crystallizes easier when the polymer chains are shorter, which was confirmed by optical polarizing microscopy. Further characterizations by temperature-modulated DSC were performed to determine whether M_n affects the specific heat capacities at the glass transition and how the rigid amorphous fraction evolves with M_n .

Choix de session parallèle

4.2 Physique des polymères: de la molécule au matériau

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Session Classification: Séance Parallèle