

Developing a low temperature spinning process for polyhydroxyalkanoates

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Polyhydroxyalkanoates (PHAs), known as bacterial polyesters, are considered novel polymers because of their biodegradability. A wide range of hydroxyalkanoate units, such as butyrates and valerates, are produced by bacterial synthesis. PHAs can be polymerized and copolymerized with varying mechanical and structural properties. Due to their biocompatibility, PHAs have been introduced in the fabrication of medical products, such as sutures and wound dressings. Some studies have explored the use of bacterial polyester for controlled release applications with thermally sensitive chemicals and drugs. Since PHAs are typically processed at temperatures as high as 200 °C, this requires a post spinning stage for chemical and drug incorporation. Hence, the development of a low temperature spinning process for bacterial polyester to prevent drawbacks of post-spinning drug incorporation, such as drug absorption that leads to an uneven release profile. To achieve this goal, we analyzed PHA solubility properties to develop a spinning process at low temperature. Next we compared dissolution of poly(3-hydroxybutyrate-4-hydroxybutyrate) (P34HB) in various solvents such as tetrahydrofuran, dioxane, methylene dichloride, and chloroform. This solvent study found methylene dichloride as the most suitable solvent. As a result, polymer solutions of various concentrations were coagulated and regenerated in methanol at different temperatures to determine the optimal coagulating conditions. The polymer films were tested for their thermal properties, molecular weight distribution and degradation profile. It was determined that the process used did not cause significant degradation in the polymer. Currently we are working on translating this process of making bacterial polyester films at low temperature to produce continuous filaments at low temperature. The project would further involve testing the effect of drug incorporation by incorporation of drugs during spinning and determining a release profile for those drugs. This study would help in developing a single step process for drug incorporation during fiber spinning, which can be utilized for drug delivery applications.

Biography

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