

Production and Characterization of Polyhydroxybutyrate Biopolymer from *Azohydromonas australica* Using Sucrose as a Sole Carbon Source

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Abstract

A bacterial strain *Azohydromonas australica* DSM 1124 has been chosen, which is accumulated intracellular poly- β -hydroxybutyrate particles, each method which has been merit and demerit for employed in PHB extraction. Our study, we have selected chloroform-sodium hypochlorite method. It is one of the methods for extracting PHB forms *Azohydromonas australica* DSM 1124. Polyhydroxybutyrate is a biodegradable and biocompatible thermoplastic with many interesting applications in medicine, food packaging and tissue engineering materials. That does not produce any toxins or residues in the environment like petroleum based plastics. The present study was emphasized on enhanced production of PHB by *Azohydromonas australica* using sucrose as a sole carbon source and estimation of biomass and sucrose content in the media. The batch kinetics analysis of *Azohydromonas australica* was done an interval of 3 h. Batch cultivation with optimized media recipe in a 7 L bioreactor exhibited a maximum biomass 1.71 g/L and PHB concentration 2.67 g/L for *A. australica*. The characterization of PHB was done by growth kinetics studies, UV-Spectrophotometer and Fourier transforms infrared spectroscopy (FTIR).

Keywords: Polyhydroxybutyrate; *Azohydromonas australica*; Sucrose; FT-IR

Introduction

Biopolymers are polymers that are generated from renewable natural sources, are often biodegradable and nontoxic. They can be produced by biological systems (like, microorganisms, plants and animals) or chemically synthesized from biological materials (like, sugars, starch, natural fats and oils). There are two strategies applied for converting these raw materials into biodegradable polymers: extraction of the native polymer from a plant or animal tissue and a chemical or biotechnological route of monomer polymerization. Biodegradable biopolymers (BDP) are an alternative way for the production of petroleum-based polymers like, traditional plastics. Some BDP degrade in only a few weeks, while the others take several months for the degradation. In principle, the properties relevant for application as well as biodegradability are determined on the basis of the molecular structure and characterization. According to the *American Society for Testing and Materials*, biopolymers are degradable in results from the action of naturally occurring microorganisms such as bacteria, fungi and algae [1]. The BDP can be produced from natural occurring raw materials such as starch, sugar, cellulose as well as fossil oils. Besides the BDP biodegradability, other relevant aspects for processing are also important including, thermal stability and viscosity that are used as a conventional technology without any large adaptations. This economic value of renewable raw materials will increase in a significant extent and stimulate new industrial activities [2,3]. The potential for biodegradable of polymers are depends on legislation, local directives and sufficient waste treatment capacity at composting plants (for packaging, disposables, foils for agricultural use). Similarly, the development in the chemical sector (like, solvents, glues, coatings) are also become an important encouragement for the application of degradable raw materials. A new biotechnological process have direct implications for society but there are significant ethical issues publicly and personally concern to sustain for the social development [4,5]. To overcome of these issues, they can widely use of BDP to prevent or reduce toxic effect of industrial waste in the environment and providing a high level of

environmental protection. The BDP also protects the loss of soil fertility and significantly enhance soil carbon level.

Biodegradable plastics are either partly or fully degraded either by non-enzymatic or enzymes activity possessed by some microorganisms that lead to a chain breakage of the polymer into monomers. Some type of the biodegradable plastics being produced polyhydroxyalkanoates (PHAs), polylactides, aliphatic polyesters, polysaccharides, and copolymers and blends of starch and polypropylene [6,7]. Blends of starch and polypropylene can be regarded only as semi-biodegradable while PHA is 100% biodegradable [8]. One of most widely used biodegradable plastics is polylactic acid (PLA) for petroleum derived plastics but it is costly than polyhydroxybutyrate (PHB) [9]. But there is several demerits in PLA including, slow crystallization, poor heat resistance and mechanical brittleness [10]. In addition, although lactic acid can currently be produced by industrial scale fermentation, it still needs chemical polymerization to get PLA, which makes the production process more complicated and time-consuming [9]. Hence, further exploration of other biodegradable plastics like PHB is warranted. PHB belongs to the class of biodegradable plastics, polyhydroxyalkanoates (PHAs).

PHB is a natural biopolymers, that are synthesized and catabolized by microorganisms particularly bacteria [11,12]. There are more than 300 of such species known to produce PHB's [13]. A number of bacteria such as *Azotobacter*, *Bacillus*, *Azohydromonas australica*,

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Archaeobacteria, Methylobacteria and Pseudomonas have been found to accumulate PHB to varying levels. *Ralstonia eutropha* (formerly *Alcaligenes eutrophus*) was found to accumulate PHBs up to 80 percent dry weight [14]. Macrae and Wilkinson [15] reported that Properties of biodegradability of PHB produced by *B. cereus* and *B. megaterium*. The shape, size, structure and physical properties of these species differ from organism to organism [16]. Beijerinck was the first to observe lucent granules of PHB in bacterial cells in 1888 [17]. Hence, in the previous studies showed that *Bacillus cereus*, *Bacillus subtilis*, *B. megaterium* have been shown to produce higher amounts of PHB using agro-industrial waste with variable level of carbon sources [18]. Besides recent studies revealed that the highest production of PHB using *Bacillus thuringiensis* along with H₂ gas, glucose: NaNO₃ in 4:1 ratio [19,20]. PHB are comparable to commonly used in bulk plastics like, polypropylene. It is a unique natural biopolymer, which have three significant features like, thermoplastic processability, 100% water and moisture resistance and 100% biodegradability [21]. It could therefore be used for applications similar to those of common plastics and would fit well into new waste-management strategies. PHB also has some addition advantages that monomer 3-hydroxybutyric acid is a chiral molecule. These monomer units of PHB can be used as basic molecules for the production in pharmaceutical or agro-chemical agents [22]. PHB is biocompatible and nontoxic in nature. The rate of PHB degradation is very slow *in vivo* and rate limiting release of drugs [23]. The aim of our study is to evaluate the biopolymer PHB producing *Azohydromonas australica* DSM 1124 from German collection of microorganisms and cell cultures (DSMZ, Germany) was obtained as a gift strain from IIT Delhi. The isolated bacterial cells were identified and characterized by morphological, biochemical, physiological and molecular characteristics. The PHB was extracted and purified after mass culture of the selected bacterial strain *Azohydromonas australica*. The purified PHB was then identified and characterized by various analytical methods such as growth kinetics study, Ultraviolet-visible Spectroscopy and Fourier Transform Infrared Spectroscopy (FTIR).

Materials and Methods

Micro-organism and maintenance

Azohydromonas australica DSM 1124 from German Collection of Microorganisms and Cell Cultures (DSMZ, Germany) was obtained as a gift strain from IIT Delhi. The strain was maintained on nutrient agar (Hi-Media, India) slants at 4°C and sub-cultured monthly.

Growth kinetics studies

Media composition for growth kinetics study of *Azohydromonas australica* was prepared containing : sucrose 25 g/L, (NH₄)₂(NO₃) 2.8 g/L, KH₂PO₄ 3.25 g/L, K₂HPO₄ 3.25 g/L, MgSO₄ 0.2 g/L (Hi-media) and trace element solution 1.5 mL/L. KH₂PO₄ and MgSO₄ were sterilized separately to prevent precipitation and added to the medium in laminar air flow. TES was filter sterilized and added to the medium in sterile environment. The parent culture of *A. australica* was inoculated (5%) in 100 ml media in different flasks. The flasks were kept in an incubator shaker at 33°C and 200 rpm. Temperature and pH was maintained at 33°C and 7, respectively. Samples were taken at intervals of 3 h for the purpose of taking optical density. Wet weight of pellets was taken to estimate biomass of organism.

Extraction of polyhydroxybutyrate (PHB) and quantification

The extraction of polyhydroxybutyrate (PHB) from *Azohydromonas australica* was described in detail by Xu and Chen [24]. Briefly, cultures were incubated for 48 h at the optimal growth temperature. Bacterial

cells were centrifuged at 6000 rpm for 15 min and were dried at room temperature for 24 h. The dry weight of the pellets was determined. Bacterial cell walls were lysed by adding sodium hypochloride, mixing and incubating at 37°C for 1 h. Supernatant was obtained by centrifugation and transferred to a Soxhlet system. Cell lipids and other molecules were treated with 5 ml of 96% ethanol and acetone. PHB was extracted by hot chloroform (adding 10 ml chloroform in a water bath). Then chloroform was evaporated to obtain PHB crystals. By adding 10 ml of 98% sulfuric acid at 60°C for 1 h in water bath after this PHB crystals were converted into crotonic acid. The absorbance of the solution was measured at 235 nm in a UV-visible spectrophotometer against a sulfuric acid used as a blank. The amount of PHB per gram dry weight of bacterial cells was determined using a standard curve of PHB.

Characterization of polyhydroxybutyrate

The polyhydroxybutyrate (PHB) was further characterized by UV-visible Spectrophotometer and Fourier Transform Infrared Spectroscopy (FTIR).

UV-visible spectrophotometer

Absorbance spectra of PHB isolated from *Azohydromonas Australica* was measured at 235 nm using sulphuric acid as a blank under UV-visible Spectrophotometer (Shimadzu, model UV 1700).

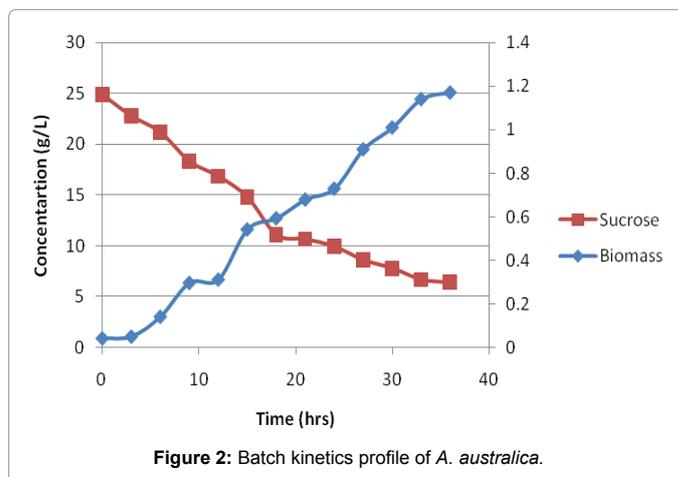
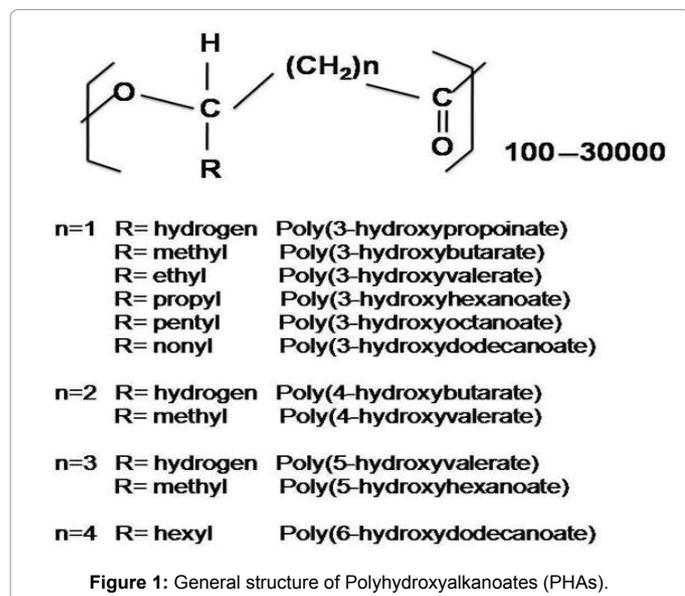
Fourier transforms infrared spectroscopy (FTIR) studies

For FTIR analyses, the PHB samples were dissolved in chloroform and then added to KBr pellets. After complete solvent evaporation, FTIR spectra were recorded using a spectrometer (Shimadzu FTIR 8201-PC). A total of 20 scans were recorded at a 2 cm⁻¹ resolution, between 4000 and 1000 cm⁻¹.

Results and Discussion

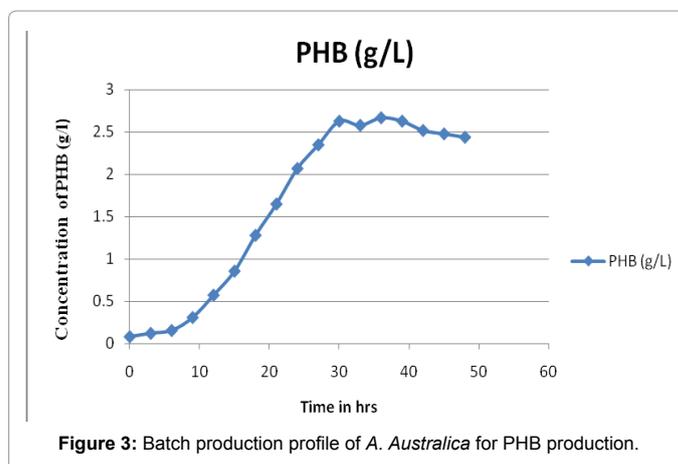
Extraction of polyhydroxybutyrate (PHB)

There are no reports on extraction of PHB from *Azohydromonas australica* spp. In our studies, we have performed only one method to extract PHB from *Azohydromonas australica* DSM 1124. Chloroform-sodium hypochlorite method is one of the PHB extractions from *Azohydromonas australica* DSM 1124, which has provided a 73% extraction rate with more than 90% purity and nontoxic for the macromolecules. After treated by sodium hypochlorite, cells break up and PHB disperse in solution. PHB were rapidly transferred to the chloroform phase, which accelerate the extraction and shorten the contact time of PHB with the sodium hypochlorite. Another biological method for the PHB production by recombinant *Bacillus subtilis* PT5(ReA)/pBmRBC was induced by IPTG. The effect of IPTG concentration and cell density on PHB yield was evaluated. PHB content increased as a function of IPTG dosage. *B. subtilis* is a GRAS organism that has many advantages for the large-scale production of chemicals, enzymes, and amino acids, including the fact that it does not produce enzymes that degrade PHA. We therefore examined in the present study whether *B. subtilis* can be an effective vehicle for PHB production (Figure 1). The PHB synthesis genes and mechanisms of *B. megaterium* and *R. eutropha* have been well characterized and served as a basis for creating a recombinant *B. subtilis* strain that can synthesize PHB than other carbon sources (Table 1). A comparative chart between the productions of PHB using *Alcaligenes eutrophus* vs. *Azohydromonas australica* as summarized in Table 1.



| S. No. | <i>Alcaligenes eutrophus</i> | <i>Azohydromonas australica</i> |
|--------|---|---|
| 1 | Glucose as a carbon source | Sucrose as a carbon source |
| 2 | PHB production equal to 80% of their dry body weight | PHB production more than 80% of their dry body weight |
| 3 | PHB obtained 15 g/L | PHB obtained 3.17 g/L |
| 4 | PHB contains 5.3% | PHB contains 29% |
| 5 | Biomass contains 36% | Biomass contains 37% |
| 6 | Maximum specific growth rate 0.075 to 0.300 h ⁻¹ | Maximum specific growth rate 0.26 h ⁻¹ |

Table 1: Comparative chart between the productions of PHB using *Alcaligenes eutrophus* vs. *Azohydromonas australica*.



Batch kinetics profile of *Azohydromonas australica*

Kinetic analysis of *Azohydromonas australica* was performed after an interval of 3 h. Biomass and sucrose content in the media was estimated by wet weight basis. Sucrose content was estimated by dinitrosalicylic (DNS) method. It was clear from the data that the exponential phase of *A. australica* begins from 9 h till 36 h after that decline phase started. The maximum biomass of 1.17 g/L was observed at 36 h and after that growth curve was declined (Figures 2 and 3 and Tables 2 and 3). Similarly, sucrose content showed significantly decreases with time as summarized in Table 2 and Figure 2.

Polyhydroxybutyrate (PHB) quantification

The estimation of PHB isolated from *A. australica* was calculated by UV-visible spectrophotometer at 235 nm as summarized in Table 3 and Figure 3. It is clear from the data that the maximum PHB concentration reaches to 2.67 g/L in 36 h (Table 3).

Characterization of the intracellular PHB

The purified PHB was characterized by UV-Spectrophotometer analysis and Fourier Transform Infrared Spectroscopy (FTIR).

Ultraviolet absorption spectroscopy

When PHB dissolved in sulphuric acid, it converted into crotonic acid, which has a maximal absorption at 235 nm. The UV spectra of samples isolated from *Azohydromonas australica* DSM 1124 that of the

| S. No. | Time | Biomass g/L | Sucrose g/L |
|--------|------|-------------|-------------|
| 1 | 0 h | 0.044 | 24.856 |
| 2 | 3 h | 0.052 | 22.764 |
| 3 | 6 h | 0.143 | 21.168 |
| 4 | 9 h | 0.298 | 18.309 |
| 5 | 12 h | 0.313 | 16.837 |
| 6 | 15 h | 0.543 | 14.778 |
| 7 | 18 h | 0.594 | 11.091 |
| 8 | 21 h | 0.68 | 10.665 |
| 9 | 24 h | 0.73 | 9.932 |
| 10 | 27 h | 0.91 | 8.605 |
| 11 | 30 h | 1.01 | 7.803 |
| 12 | 33 h | 1.14 | 6.687 |
| 13 | 36 h | 1.17 | 6.434 |
| 14 | 39 h | 1.03 | 6.312 |
| 15 | 42 h | 0.98 | 5.567 |
| 16 | 45 h | 0.95 | 5.333 |
| 17 | 48 h | 0.84 | 5.189 |

Table 2: Batch kinetics of *A. australica* for biomass and sucrose concentration.

standard completely. According to the peak value, the content of PHB can be calculated by a PHB standard curve (data not shown) [25-28]. Concentrated H₂SO₄ destroys cell membrane, and the fragments of cell have no absorption at 235 nm. Therefore, the quantity of PHB in ferment medium is calculated by digesting the cells directly.

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