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Research Article

Comparative Account on Proliferation Rate of Microalgae Used in Biodiesel Production by Indigenously Prepared Bioreactors

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Abstract: Algae as a feedstock is emerging at the forefront of biofuel research due to increasing awareness of global energy issues in conjunction with the production limitation of agriculture based oilseed crops. In this concern, India is lagging behind with respect to research emphasis. The long term Research and development strategies were implemented in the developing countries for biofuels from different sources. One of the most fascinating areas of researcher always remained with algae as a source of biofuel. All the above the more research than development will be required in this field. This paper focuses the comparative Proliferation rate of Chlorophycean algae in indigenously prepared bioreactors as well as drinking bottled models.

Keywords: Indigenously prepared bioreactor, Biofuels, Algae.

1. Introduction

Microalgae are single-cell, photosynthetic organisms known for their rapid growth and highenergy content. Some algal strains are capable of doubling their mass several times per day. In some cases, more than half of that mass consists of lipids or triacylglycerides-the same material found in vegetable oils (9). These bio-oils can be used to produce such advanced biofuels as biodiesel, green diesel, green gasoline, and green jet fuel. Higher oil prices and increased interest in energy security have stimulated new public and private investment in algal biofuels research (8). Though the Chlorophycean microalgae owe the higher concentration of oil content they are concealed because of cultivation strategies need to use and high economic investments required for engineering and construction of basic infrastructure. This is due in time that the high cost of even simple algae production system and algal mass culture technology, maintenance of algal strains in the

cultivation system for achievement of high productivities of biomass with a high content of lipids or other biofuel precursors (11).

Algae range from small, single-celled organisms to multi-cellular organisms, some with fairly complex and differentiated form. Like plants, algae require primarily three components to grow: sunlight, carbon-dioxide and water. Photosynthesis is an important biochemical process in which plants, algae, and some bacteria convert the energy of sunlight to chemical energy. Microalgae contain lipids and fatty acids as membrane components, storage products, metabolites and sources of energy (2).

Algal oil processes into biodiesel as easily as oil derived from land-based crops. The difficulties in efficient biodiesel production from algae lie not in the extraction of the oil, but in finding an algal strain with a high lipid content and fast growth rate that isn't too difficult to harvest, and a cost-effective cultivation system (i.e. type of photobioreactor) that is best suited to that strain (1). Microalgae have much faster growth rates than terrestrial crops. The per unit area yield of oil from algae is estimated to be from between 5,000 to 20,000 gallons (18,927 to 75,708 litres) per acre, per year; this is 7 to 31 times greater than the next best crop, palm oil (635 gallons or 2,404 litres) (11).

The production of algae to harvest oil for biodiesel has not yet been undertaken on a commercial scale, but feasibility studies have been conducted to arrive at the above yield estimate. In addition to its projected high yield, algaculture — unlike crop-based biofuels do not entail a decrease in food production, since it requires neither farmland. Many companies are pursuing the development of algae bioreactors for various purposes – including biodiesel production and CO₂ capturing (10).

Producing biodiesel from algae has been touted as the most efficient way to make biodiesel fuel. The main advantages of deriving biodiesel from algae oil include: rapid growth rates, a high per acre yield (7), algae biofuel contains no sulphur, algae biofuel is non-toxic, algae biofuel is highly biodegradable, and algae consume carbon dioxide as they grow, so they could be used to capture CO_2 from power stations and other industrial plant that would otherwise go into the atmosphere (4).

Research into algae for the mass production of oil is mainly focused on microalgae. The preference towards microalgae is due largely to its less complex structure, fast growth rate, and high oil content. Attempts were made in this paper to show the rate of multiplication of two microalgae, *Chlorella vulgaris* and *Scenedesmus dimorphus*.

2. Material and method

2.1 Collection of algal sample

The Samples of microalgae were collected from ponds, puddles, dams and rivers of Ahmednagar district with the help of plankton net made up of bolting silk (No. 25; mesh size 55 micron). The samples were then transferred to narrow mouthed plastic bottles and brought to the laboratory.

2.2 Isolation of algae

The collected samples were brought to the laboratory and isolated by the serial dilution method followed by plating. Media used for isolation and to maintain the culture were Bold Basal media, modified Chu 13 media (5) and incubated at $25 \pm 1^{\circ}$ C under 1.2 ± 0.2 Klux light intensity with 16:8 hrs light photoperiod.

2.3 Studies of multiplication rate of microalgae

The growth rate of both algae *Chlorella vulgaris* and *Dunaliella salina* were studied by growing algae in Erlenmeyer flask (250ml), indigenously prepared photobioreactor with the help of cleaned electricity tube bars and empty drinking water bottles (1 liter). As shown in photo plate 1 incubated at $25 \pm 1^{\circ}$ C under 1.2 ± 0.2 Klux light intensity with 16:8 hrs light photoperiod. Artificial air was provided with the help of bower for the indigenously prepared model. The experiment was conducted in triplicates.

2.4 Measurement of growth of the algae

The rate of growth was calculated after 15 days with the help of proliferation rate (Huang *et al.*, 2002):

$$K = \frac{(logOD_t - logOD_0)}{T} X 3.322$$

Where, OD_i : Terminal optical density; OD_0 : Initial optical density; T: Days Optical density was measured by spectrophotometer using 680nm wavelengths (6) each sample was thoroughly swirled before taking the measurement. The calculated value was recorded in Table 1.

3. Results

As shown in Table 1 the proliferation rate of Chlorella and Scenedesmus had shown significant values in indigenously prepared photobioreactor and bottled bioreactors. In Erlenmeyer flask, the proliferation rate of Chlorella was 0.031 which is not significant whereas it shows significant values in indigenously prepared electricity tube bioreactors, 0.098 and water bottle bioreactor 0.176. Same were observations with Scenedesmus it had a non-significant proliferation rate in Erlenmeyer flask i.e. 0.042, whereas significant in indigenously prepared electricity tube bioreactors, 0.0112 and water bottle bioreactor 0.190. In comparison with the proliferation rate of both algae, Scenedesmus shows more significant growth over Chlorella at culture condition in modified Chu 13 liquid medium. Apart from this, the Chu 13 medium proved good for the growth and isolation of the unicellular Chlorophycean algae.

Table 1. Effect of different cultivation conditions for biomass multiplication on the growth of algae (Proliferation rate).

| Sr. No. | Name of algae | Erlenmeyer flask | Indigenous Bioreactor | Drinking water Bottles |
|---------|-----------------------|---------------------------|-----------------------|-------------------------------|
| | | Rate of proliferation (K) | | |
| 1 | Chlorella vulgaris | 0.031 | 0.098* | 0.176* |
| 2 | Scenedesmus dimorphus | 0.042 | 0.112* | 0.190* |

*Significance observed on the proliferation rate in different growth condition ($p \le 0.05$).

Figure-1- Comparison of proliferation rate of Chlorella and Scenedesmus by Indigenously prepared bioreactore models



^{4.} Discussion

The growth rate observed in the different models will prove a concrete step towards the production and increase in the yield of biomass for oil production. The proliferation rate shown by both algae in the bioreactor was due to availability of space and facility of aeration to the algae. In the bioreactor prepared from the drinking water exchange of gas was easier as compared to tube bioreactor. Also, the byproduct of photosynthesis i.e. oxygen gets escaped easily from this bioreactor. Availability of CO₂ was easy due to continuous circulation of air through media. Assuming the technology can be developed, become worthwhile; perhaps even a significant, contribution to the goal of renewable energy production, in particular of transportation fuels. Again, the resource potential requires further studies. The different algae should be checked for the maximum proliferation rate and standardized.

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