Optimization production conditions of photosynthetic purple bacteria biomass at pilot scale to remove sulphide from aquaculture pond

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For the purpose of sulphide removal in aquaculture ponds, three strains (name: TH21, QN71, QN51) were isolated and selected with the highest sulphide removal activity from Thanh Hoa and Quang Ninh coastal zones. These strains have been identified and tested in a number of aquaculture ponds in different areas with good water quality results. With the objective of purple non sulfur bacteria biomass production containing 3 selected strains for wide application and suitable price for farmers, in this study, we study on optimum conditions of mixed purple non sulfur bacteria biomass production at pilot scale. The results showed that the sources of substrates were soybean meal (1g/l) and acetate (0.5g/l). These substrates are low cost, easy to find, convenient in large culture. The mixture of photosynthetic bacteria can be cultured in glass tanks, under micro aerobic and natural lighting conditions that produce highly concentrated photosynthetic bacteria and lowest rest media.

Keywords: purple non sulfur bacteria, Rhodovulum sulfidophilum; Rhodobacter sphaeroides; sulphide

1. Introduction

Sulphide is generated from the rest feed and animal waste, which accumulates at the bottom of the aquaculture ponds and is converted an aerobically by microbes (Kutako et al., 2009). The toxicity of sulphide is mainly due to the inhibition of cytochrome C oxidase and to interference with other crucial enzymes, mainly by H2S (Affonso et al., 2002; Konishi et al., 2013). The presence of sulphide also can provide a favourable condition for bacterial proliferation and increase the rate of bacterial infection in sediment-dwelling animals (Bourgeois and Felder, 2001). The studies of sulphide on aquatic animals indicate that...
2. Materials and methods

2.1. Materials

The study was carried out by using mixture of purple non-sulphur bacteria strains including *Rhodobacter sphaeroides* QN71, *Rhodobacter sphaeroides* QN82 and *Rhodovulum sulfidophilum* TH3 with high sulphide removal capacity. Three strains were isolated from the bottom of aquaculture ponds from Thanh Hoa and Quang Ninh coastal. A number of biological characteristics were investigated at Environmental Biotechnology Laboratory-Institute of Biotechnology, Vietnam Academy of Science and Technology.

DSMZ 27 medium which consists of the following components: (1L); K₂HPO₄: 1 g; KH₂PO₄; 0.5 g; MgSO₄; 7H₂O; 0.4 g; NaCl; 15 g; NH₄Cl: 0.4 g; CaCl₂ 2H₂O: 0.05 g; yeast extract: 0.3 g; sodium acetate: 1 g; succinic acid: 0.5 g; trace element solution: 1 ml; vitamin solution: 1ml; agar: 20 g; distilled water to 1000 ml; pH of the medium: 6.8–7.0 before autoclaving. The trace element solution (1L): HCl (25%) 6.5 ml; FeCl₃·4H₂O 1.5 g; H₂BO₃ 0.3 g; MnCl₂·2H₂O 0.03 g; CoCl₂·6H₂O 0.2 g; ZnSO₄·7H₂O 0.1 g; CuCl₂·2H₂O 17 mg; NiCl₂·6H₂O 24 mg; Na₂MoO₄·2H₂O 36 mg, H₂O 993 ml: 1 ml. The vitamin solution (1L) contain thiamine, 500 µg; niacin, 500 µg and biotin, 15 µg in 1000 ml of distilled water is disinfected with a filter and added to the medium before use.

2.2. Methods

Culturing mixture of three purple non sulfur bacteria

In the laboratory: mixture of purple non sulfur bacteria are cultured in glass flask or cylindrical glass jars with a rubber cap. The medium in the glass flasks is aerated with sterilized nitrogen gas replacing oxygen in the medium then dissolved oxygen concentration is 0 mg/l. Mixture of purple non sulphur bacteria are cultured under anaerobic conditions, and their growth in the logarithmic phase with a cell density of about 10⁸ transferred to a test flask with a volume of about 5-10% (v/v) to reach an initial density at OD800 of 0.1. The light intensity at the surface of the tests were controlled by incandescent lamps

Pilot scale: mixed purple non sulphur bacteria are cultured in plastic tanks or glass tanks with volume of 100 litter containing soybean and acetate media, placed outdoors in natural light.

Growth of purple photosynthetic bacteria assessment

Evaluating the growth capacity of the bacteria by determining the optical density increase of the cell suspension at the 800nm wavelength (OD₈₀₀) on the spectrophotometer.

Determination of COD content: Mixture of purple non sulphur bacteria were centrifuged at 9000 r/min for 10 min. The supernatant was used to analysis COD by APHA standard methods [3].

Statistical analysis: All experiments in this study were conducted in three replicates. Mean values and the standard deviations are presented. Analysis of the data using one-way ANOVA and significant differences at a p-value <0.05.
3. Results and discussion

3.1. Selection of some suitable substrate sources of culturing media

Although the mixture of three purple non sulfur bacterial strains were well grown in the DSMZ 27 media, it is aimed to select the appropriate substrates which are easy to use and cost effective for biomass production at pilot scale. We have determined the biomass accumulation capacity of the mixed purple non sulfur bacterial strains in DSMZ 27 media, carbon sources in media were replaced by some kinds of substrates such as acetate, succinate, glucose, glutamate, rice flour or soybean powder with amount of 1 g/l. The experiment was conducted under anaerobic conditions. The results determining the biomass accumulation of the mixture purple non sulfur bacterial strains for 5 days were estimated by ΔOD800 and are presented in Figure 1 and Figure 2.

The results showed that the mixture grow well on media containing substrates such as acetate, succinate, glucose, glutamate, rice flour and soybean powder. In particular, the growth of strains is the best on the medium containing the substrate of soybean powder. However, the DSMZ 27 media were replaced carbon by soybean powder, which include many expensive chemicals such as yeast, trace element and contain many components. It is not convenient to produce biomass at pilot scale. Therefore, we continue to test with media that contain only substrate sources of acetate, succinate, glucose, glutamate, rice flour and soybean powder with 1 g/l. The experiment was preceded under anaerobic conditions. The results of determining the biomass accumulation (in accordance with ΔOD800) of the mixed strains of purple non sulfur bacteria for 5 days are presented in Figure 3 and 4.

The results indicated that the mixture of purple non sulphur bacterial strains grew best on media containing only soybean powder with significant difference from each other media containing acetate, succinate, glucose, glutamate or rice flour (p≤0.05). Hence, soybean powder was used for further experiment.

3.2. Selection of the optimal soybean powder concentrations for biomass production

Bioproduct of purple non sulphur bacteria will be added directly to the aquaculture ponds periodically to treat sulphide, so bioproduct of purple non sulphur bacteria with the rest nutrient will pollute the secondary environment. Thus, study to find the exact composition of media which is just enough for the high-density growth of purple non sulphur bacteria and no nutrient redundancy is necessary.

In this experiment, mixture of three purple bacterial strains are cultured in media containing soybean powder at concentrations from 0.2g/l to 2g/l. Results of their biomass accumulation (by ΔOD800) after 5 culture days are shown in Figure 5 and 6.
Figure 5. Levels of biomass accumulation in different soybean concentrations

Figure 6. The mixed were cultured on different concentrations of soybean media

The results indicated that the density of mixed purple non sulphur bacteria was increased along with increasing of soybean concentration. The higher concentration, the greater density of mixed purple non sulphur bacteria in the medium. There was a gradual increase in biomass from the 0.2g/L concentration to the 2g/l concentration. At a substrate concentration of less than 1 g/l (0.2 and 0.5 g/l), the growth of strains of purple non sulphur bacteria is poorly, ranging from 28.5 to 72.6%. Collected biomass in media containing soybean concentrations from 1 to 2 g/l is higher than in the DSMZ 27 media. The difference in biomass accumulation at concentrations of 1 with 1.5, 1.7 and 2 g/l is no statistically significant difference (p ≥ 0.05).

We conducted an assessment of excess nutrient concentrations in the culture medium (by COD) at soybean concentrations from 1-2g/l. Result are shown in Table 1.

Table 1. Remaining COD concentration in culture media

| Soybean concentrations (g/l) | COD (O2mg/l) |
|-----------------------------|--------------|
| 1                           | 92           |
| 1.5                         | 247          |
| 1.7                         | 535          |
| 2                           | 864          |

The results showed that the highest biomass production was achieved in medium containing soybean powder (1g/l) supplemented with acetate carbon (at 0.5 and 1 g/l concentrations). The biomass accumulation in these media and soybean medium containing 2 g/l soybean meal were not statistically significant different (p≥0.05). The COD concentration in the medium containing soybean (1g/l) and acetate (0.5g/l) was the lowest. Thus, it is possible to use medium containing soybean (1g/l) and acetate (0.5g/l) to produce purple nonsulfur bacteria biomass at pilot scale for high biomass accumulation and the lowest rest media.

3.3. Study on combining soybean powder with other carbon sources

The objective of this part is to improve the substrate quality and expand the production of purple non sulfur bacteria without leaving the high rate of COD. Acetate and glutamate were carbon sources which are low cost, easy to find and quite good for purple non sulfur bacteria growth (Fig 4). Culturing medium contains soybean (1 g/l) supplemented with another carbon sources such as acetate or glutamate (concentrations from 0.5 – 1 g/l).

After five days of standing culture in illumination, the results were presented on the Table 2.

Table 2. Growth and remaining COD in the culture media

| Media                  | ΔOD(800) | COD (O2mg/l) |
|------------------------|----------|--------------|
| Soybean (1 g/l) add 0.5 g/l carbon sources | Acetate 1,515±0.05 | 74 |
| Soybean (1 g/l) add 1 g/l carbon sources | Acetate 1,530±0.05 | 97 |
| Soybean (2 g/l)        | Glutamate 1,432±0.04 | 117 |
| DSMZ - 27              |           | 1,205±0.02 |

The results in Table 2 showed that the highest biomass accumulation was in the medium containing soybean powder (1 g/l) supplemented with carbon sources (at 0.5 and 1 g/l concentrations). The biomass accumulation in these media and soybean medium containing 2 g/l soybean meal was not statistically significant different (p≥0.05). The COD concentration in the medium containing soybean (1g/l) and acetate (0.5g/l) was the lowest. Thus, it is possible to use medium containing soybean (1g/l) and acetate (0.5g/l) to produce purple nonsulfur bacteria biomass at pilot scale for high biomass accumulation and the lowest rest media.

3.4. Study on the effect of culture conditions on purple bacteria biomass production

Purple non sulfur bacteria grow optimally on anaerobic conditions with illuminate, but for pilot biomass production, anaerobic conditions such as in the laboratory are difficult. So, we conducted the culture in media containing soybean meal (1g/l) and acetate (0.5g/l) in three conditions: anaerobic (nitrogen aeration, rubber cap, oxygen concentration ¥ 0mg / l); micro aerobic condition (static flask, oxygen concentration from 0.5 -1mg/l) and aerobic (shake 120 rpm, oxygen concentration> 6 mg/l) under light. Results of their biomass accumulation (by ΔOD800) after 5 culture days are shown in Figure 7.
growth of the mixed was lowest, there was statistically significant difference compared with anaerobic and micro aerobic conditions. Thus, mixed purple non sulfur bacteria are not only the ability to grow in anaerobic conditions - light, but also in micro aerobic conditions - light. Therefore, the production of biomass in large tanks does not require absolute anaerobic.

3.5. Study the growth of purple photosynthetic bacteria on different material containers to produce biomass at pilot scale

Figure 8. PNSB culture in different material bioreactors

After finding the appropriate substrate sources and concentrations, we conducted PNSB culture at pilot scale, such as glass tanks (100 l) and plastic tanks (100 l), under light-anaerobic condition. The aim of this experiment is finding the material that make the best condition for the biomass production of PNSB. The experiment was conducted in natural conditions, with a temperature of about 35-37°C, biomass accumulation was monitored daily. Results were shown in Figure 8.

The results showed that the growth of PSNB in natural conditions affected by the day-night cycle and by weather, the growth of purple non sulphur bacteria was slower than conditions of light in laboratory. The growth rate in glass tank was faster than plastic tank. It takes about 10 days for high density in glass tank and 15 days in plastic tank. This explains that glass tanks can absorb more light than plastic tanks. Thus, production of biomass in scale pilot, we can use glass tanks to culture the mixed purple non sulphur bacteria to make bio-product.

4. Conclusion

- The medium for culturing of mixed purple non sulphur bacteria on pilot scale contains soybean powder (1g/l) and acetate (0.5g/l). Medium is cost effective, high biomass accumulation and no surplus substance.
- The glass tank was selected to culture purple non sulphur bacteria to obtain high biomass at pilot scale.

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5. References

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