OPTIMIZATION OF RIFAMYCIN B USING NOCARDIA MEDITERRANE A (ATCC 13865) BY FED BATCH MODE IN SHAKE FLASKS

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ABSTRACT

The present study was carried out to evaluate the optimum Production of Rifamycin B by using fed-batch mode in the shake flask at the definite level of different agitation and aeration rate, amongst different carbohydrate in the form of monosaccharide. Amongst the various carbohydrate (Arabinose, Fructose, Galactose, Glucose, Glycerol, Maltose, Mannitol, Sorbitol, and Sucrose) sources maximum, the rifamycin (578 µg/ml) with 10% glucose level was found as compared to control which was (138 ± 2.70µgml-1). The environmental factors such as incubation temperature. The optimum temperature for maximum antibiotic production (525µgml-1) was found to be 29°C, initial pH, aeration and agitation rates were also optimized. The optimum rifamycin yield (950 to 1100µgml-1) was obtained in 1.0 L fermenter with working volume of 1.5 L at 28 °C with initial pH of 6.2 to 6.5. The agitation and aeration were 300 rpm and 0.5 LL-1min-1 respectively.

Keywords: Nocardia mediterranea ATCC 13865, Agitator, Carbohydrate, Rifamycin B, Fed-batch, Optimization.

INTRODUCTION

One of the purposes of fed-batching is to make the substrate(s) available to the microorganism at an optimum concentration for a longer period of time or at specified times of the process. In the fed-batch mode, gradual addition of concentrated additives, e.g. nutrients, takes place. Fed-batch cultures can be used for fermentation processes in which the products are either growth associated or non-growth associated. In the course of the development of the fermentation process for production of rifamycin B, fed-batch fermentations have been developed in order to extend the production phase, where sources of sugar, organic and inorganic nitrogen were fed -batch wise or continuously1. The bacterial culture of Nocardia mediterranea ATCC 13865 was used for production of ant tubercular antibiotic rifamycin by submerged fermentation procedure. In previous (1st paper reference). The culture was improved by sequential mutagenic treatment with NTG (N-methyl-N'-nitro-N-nitroso-guanidine) and U.V radiation. The mutant strain was designed as Nocardia Mediterranea M-120. Carbohydrates in the form of monosaccharide’s and disaccharides were employed for the production of rifamycin. Among all sugars glucose (9.5%w/v) gave the maximum yield of antibiotic 2,5.

Stock culture

Pure and high yielding culture of *Nocardia mediterranea* ATCC 13865 was used for the production of antibiotic, rifamycin B. The culture was maintained on ML-sporulating agar medium (mineral base lactose medium) as described by figure 1. The ingredients were dissolved in distilled water. The pH of the medium was adjusted with NaOH or HCl. All the media unless otherwise stated were autoclaved at 121°C for 15 minutes. Slants were incubated at 28°C for 5 days for sporulation and then stored in the refrigerator at 4°C for further use (O.M., 2004). Sub culturing was made after every four night. The culture was improved by sequential mutagenic treatment with nitrosoguanidine (N.T.G) and UV irradiation and the selection was based on increasing concentrations of rifamycin B in shake flasks.

Fermentation medium

The initial fermentation medium for fermentation experiments containing (g l⁻¹) Glucose- 94, Soybean meal- 10, Peanut meal- 21.4, CaCO₃- 9.5, KH₂PO₄- 0.4, MgSO₄.7H₂O- 1.0, Barbitral- 2.0, Glucose- 70, Peptone-30, Glycerol- 20, CaC₂O₄- 8.0, KH₂PO₄- 19, MgSO₄.7H₂O- 1.0, Barbitral- 2.0, (NH₄)₂S₀₄- 3.0, (NH₄)₂S₀₄- 8.0, CuS₀₄.5H₂O- 0.0165, ZnS₀₄.7H₂O- 0.004, FeS₀₄.2H₂O- 0.002, MnS₀₄.4H₂O- 0.001 .was used and various other chemical such as glucose, ammonium sulphate, KH₂PO₄ and calcium carbonate etc. were of analytical grade. Fermentation experiment were conducted in 250ml shake flask with 50ml medium for a period of 12 days by changing the various concentration of media component and optimizing the cultural condition of the strain to improve the rifamycin production.

RESULT AND DISCUSSION

Production of Rifamycin

Effect of incubation temperature

The influence of incubation temperature on rifamycin B production was investigated by cultivating *Nocardia mediterranea* in medium M-6 at different levels of temperature. Initially rifamycin production was enhanced with the rise in temperature. The optimum temperature for maximum antibiotic production was found to be 29°C. An increase or decrease further in the temperature reduced the yield of antibiotic indicating that the bacterium is sensitive to temperature in many reports the optimum temperature for rifamycin B fermentation is given as 29 °C.

![Figure 1: Effect of incubation temperature on the production of rifamycin](image)

Effect of aeration

The availability of oxygen to the microbes during fermentation plays important role for the synthesis of secondary metabolites. In order to compare the effect of aeration on the bacterial cell mass and rifamycin production, the different volumes such as 25, 50, 75, 100, and 125 ml of the medium M-6 was taken into 250 ml flask. After incubation the fermentation was carried out for 120 hours. The shake flasks used had the same size and form and shaking was done under the same conditions. The lesser the volume of the medium present in the flask, the higher was the oxygen supply to the culture. The rifamycin potency was slightly increased from 350 to 362µgml⁻¹ when volume of the medium in flasks was changed from 25ml to 50ml and percent mycelia volume (PMV) varied from 24 to 26 %. Both the rifamycin production and cell biomass adversely affected when the volume was increased beyond 50 ml/flask.

The rifamycin yield was only 191 µgml⁻¹ and cell biomass formation was poor (PMV) when 125 ml of the medium was present in the flask. The shaking of flasks on rotary mechanical shaker provided aeration for the microorganism. The rifamycin production was maximal when 50 ml of fermentation medium was present in the flask. Further increase resulted in the volume in lowering the antibiotic production due to lesser agitation, which in turn resulted in to poor oxygen supply to the culture. The optimum volume of production medium (50ml/250ml) flask is in accordance with the studies of BM.°².°¹.°.
Effect of aeration and agitation

The availability of oxygen to the microbes during fermentation is very critical for the synthesis of enzymes or metabolites. Only dissolved oxygen becomes available to the microbes for its propagation in metabolites formation. To investigate the effect of aeration on the production of antibiotic rifamycin B, a series of batch cultures were run using Nocardia mediterranea M-120. The rate of agitation was set at 200, 300 and 500 rpm with different rates aeration (1.0 to 2.0 LL·min⁻¹). During the course of fermentation, pH, residual glucose, cell biomass and antibiotic potency were estimated. In this processes the optimum rifamycin yield (949 to 1100 µg ml⁻¹) was obtained in 1.0 L fermenter with working volume of 1.5 L at 28 °C with initial pH of 6.2 to 6.5. The agitation and aeration were 300 rpm and 1.0 LL·min⁻¹ respectively.

Effect of carbohydrates

An adequate supply of the carbon energy source is critical for optimal growth and production formation. The culture Nocardia mediterranea M-120 was cultivated in the basal medium M-6 deleting glucose. The effect of different carbohydrates such as glucose, lactose, fructose, sucrose, maltose and glycerol etc on the production was investigated in the basal medium. Each carbon source was added separately at a concentration of 5%, 7.5% and 10% (w/v) to the fermentation medium before sterilization in all the experiments to investigate their effect on the rifamycin production. Glucose showed the highest production of antibiotic (578 µg ml⁻¹). Fructose also stimulated the rifamycin production to a lesser extent as compared to glucose. The antibiotic activity was maximal in the presence of 10% glucose (578 µg ml⁻¹) and 10% fructose.
(489 µg/ml). While sucrose, sorbitol, galactose and maltose were less effective.

Addition of xylose, lactose or starch even decreased the antibiotic production. The control basal medium without any sugar gave 138 µg/ml rifamycin B. It follows that defatted meals also provide nutrients both for bacterial growth and antibiotic production. The addition of sugar however, increases the synthesis of antibiotic.

![Figure 5: Effect of Aeration on the Production of Rifamycin B by Nocardia Mediterranea M-120 at 500 Rpm](image)

![Figure 6: Effect of different carbohydrates on the production of rifamycin](image)

**REFERENCES**


