## Preliminary studies on Saccharomyces cerevisiae cell mass production

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This paper describes the preliminary studies on factors that affect the growth of Saccharomyces cerevisiae strain from Fermipan. The fermentation medium and inoculum medium were the same and contained (gl<sup>-1</sup>); yeast extract, 2.5; bacterial peptone. 1.15; (NH<sub>4</sub>) 2 HPO<sub>4</sub>, 0.25 and MgSO<sub>4</sub>. 7H<sub>2</sub>O, 0.025. The fermentation was carried out at 30°C and pH 5.0, while shaking at 100rpm. The growth was monitored as dry weight. In the fermentation medium containing 50,100 and 150 gl<sup>-1</sup> glucose, dry weight of yeast obtained was 4.7, 9.5 and 9.8gf<sup>-1</sup> respectively and the residual glucose present at 24h of fermentation was 0, 12.2 and 48.4gl-1 respectively. As the residual sugar in 150gl<sup>-1</sup> glucose containing medium was 48.4gl<sup>-1</sup> it was decided to do further studies in medium containing 50 and 100 gl<sup>-1</sup> glucose. The impact of addition of different amounts of inorganic nitrogen source on production of biomass and alcohol was studied. The medium containing glucose 50gl-1 with 0.25 and 0.50 gl-1 (NH<sub>4</sub>), HPO4 along with other nutrients the dry weight of yeast obtained was 4.4 and 4.7 and the alcohol produced was 12.5 and 10.8gl<sup>-1</sup> respectively. While with 100gl<sup>-1</sup> glucose medium with 0.25 and 0.50 gl<sup>-1</sup> (NH<sub>4</sub>)<sub>2</sub> HPO<sub>4</sub> along with other nutrients, the dry weight of biomass was 9.4 and 9.9gl<sup>-1</sup> and alcohol produced was 20.5 and 18.5gl<sup>-1</sup> respectively. Complete glucose utilization was obtained in 50gl<sup>-1</sup> glucose with 0.25 and 0.5gl<sup>-1</sup> (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> containing medium. Residual sugar present in the 100gl<sup>-1</sup> glucose and 0.25 and 0.5gl<sup>-1</sup> (NH<sub>4</sub>) <sub>2</sub> HPO<sub>4</sub> containing media were 6.7 and 1.3gl<sup>-1</sup> respectively. The glucose concentration of 100gl1 was selected for further studies as the glucose in the medium containing 0.5g of (NH<sub>4</sub>) <sub>2HPO4</sub> was almost completely utilized by cells. In the next set of experiment, the medium contained 100gl<sup>-1</sup> glucose, along with 0.25, 0.50 or 0.75 gl<sup>-1</sup> (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> and other nutrients. The dry weight obtained and alcohols produced were 7.3, 8.1 and 7.5gl<sup>-1</sup> and 28.8, 25.3 and 23.0gl<sup>-1</sup> respectively. Hence the growth medium containing glucose, 100 gl<sup>-1</sup> and (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub>, 0.5gl<sup>-1</sup> along with other nutrient was selected for further studies. The effect of yeast extract on growth and In the medium containing 100gl<sup>-1</sup> glucose and alcohol fermentation was studied. 0.5 gl<sup>-1</sup> (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> supplemented with 2.5, 3.5 and 4.5gl<sup>-1</sup> yeast extract, the cell mass of 7.7, 9.1 and 10.7 gl<sup>-1</sup> and alcohol of 24, 22.5 and 20.5 gl<sup>-1</sup> were produced respectively. As yeast extract addition was economically not feasible it was decided to keep the yeast extract at 2.5gl1 level and increase the (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub> to 0.5g. With increase in (NH<sub>4</sub>)<sub>2</sub>HPO<sub>4</sub>, the pH of the medium decreased from 5.0 to 3.0 during fermentation. Hence the effect of pH on growth of the S.cerevisieae was studied. The initial pH was adjusted to 5.0, 6.0 and 6.5 and the pH was adjusted 4 hourly using 4N sterile NaOH. The dry weight obtained was 6.0, 7.0 and 6.7gl1 respectively. The alcohol produced was 30, 28 and 25gl<sup>-1</sup> respectively. Hence the pH was maintained around 6.0 during the fermentation.