

## Enhancement of $\beta$ -mannanase production by *Bacillus subtilis* ATCC11774 through optimization of medium composition

### ABSTRACT

Palm kernel cake (PKC) has been largely produced in Malaysia as one of the cheap and abundant agro-waste by-products from the palm oil industry and it contains high fiber (mannan) content. The present study aimed to produce  $\beta$ -mannanase by *Bacillus subtilis* ATCC11774 via optimization of the medium composition using palm kernel cake as substrate in semi-solid fermentation. The fermentation nutrients such as PKC, peptone, yeast extract, sodium chloride, magnesium sulphate ( $\text{MgSO}_2$ ), initial culture pH and temperature were screened using a PlackettBurman design. The three most significant factors identified, PKC, peptone and NaCl, were further optimized using central composite design (CCD), a response surface methodology (RSM) approach, where yeast extract and  $\text{MgSO}_2$  were fixed as a constant factor. The maximum  $\beta$ mannanase activity predicted by CCD under the optimum medium composition of 16.50 g/L PKC, 19.59 g/L peptone, 3.00 g/L yeast extract, 2.72 g/L NaCl and 0.2 g/L  $\text{MgSO}_2$  was 799 U/mL. The validated  $\beta$ -mannanase activity was 805.12 U/mL, which was close to the predicted  $\beta$ -mannanase activity. As a comparison, commercial media such as nutrient broth, M9 and Luria bertani were used for the production of  $\beta$ -mannanase with activities achieved at  $204.16 \pm 9.21$  U/mL, 50.32 U/mL and 88.90 U/mL, respectively. The optimized PKC fermentation medium was four times higher than nutrient broth. Hence, it could be a potential fermentation substrate for the production of  $\beta$ -mannanase activity by *Bacillus subtilis* ATCC11774.

**Keyword:**  $\beta$ -mannanase; *Bacillus subtilis*; Palm kernel cake; Optimization; Response surface methodology