

## Evaluation of the enzymatic activity of immobilized *Lysobacter sp.* microorganisms

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### INTRODUCTION & AIM

The specific enzymatic activity of bacterial cultures illustrates the key applications of a studied strain of bacteria and the biological products based on it.

The aim of this study was to determine the enzymatic activity of immobilized *Lysobacter sp.* cells on inorganic matrices.

### METHOD

The assessment of proteolytic activity was determined by the spectrophotometric method using a UV/Vis spectrophotometer for studying nano- and microvolumes of liquids NABI NICRODIGITAL. A 1% aqueous solution of casein was used as a substrate.

A unit of  $\beta$ -1,4-glucanase activity was defined as the amount of enzyme that can produce 1  $\mu$ mol of glucose in 1 minute according to the assay condition.

The Bradford method was used to determine protein concentration. Bovine serum albumin (BSA) was used as a standard. Fractions with the highest  $\beta$ 1,4-glucanase activity were pooled and lyophilized.

### RESULTS & DISCUSSION

The maximum proteolytic activity of enzymes in the culture liquid of the studied *Lysobacter* strain is observed at a temperature of 40°C.

Immobilization of cells on carriers generally has a positive effect on enzyme production and activity. In the case of immobilization on sodium carboxymethylcellulose and the sodium form of montmorillonite, the temperature range of enzymatic activity expands from 35°C to 45°C, which will expand the use of biocompositions based on immobilized cells and their enzymes.

The proteolytic activity of extracellular enzymes in the temperature range 20-40 and the activity of  $\beta$ -1,4-glucanase were studied, and the influence of pH and temperature on the activity and stability of the enzyme was revealed.

Optimal cultivation conditions for the *Lysobacter* strain were selected for the highest production of  $\beta$ -1,4-glucanase: glucose, 1% by weight; yeast extract, 0.2% by weight; CMC, 0.75% by weight; pH 7; amount of inoculum, 3.5% by volume; and optimal temperature for incubation, 30°C (Figure 1).

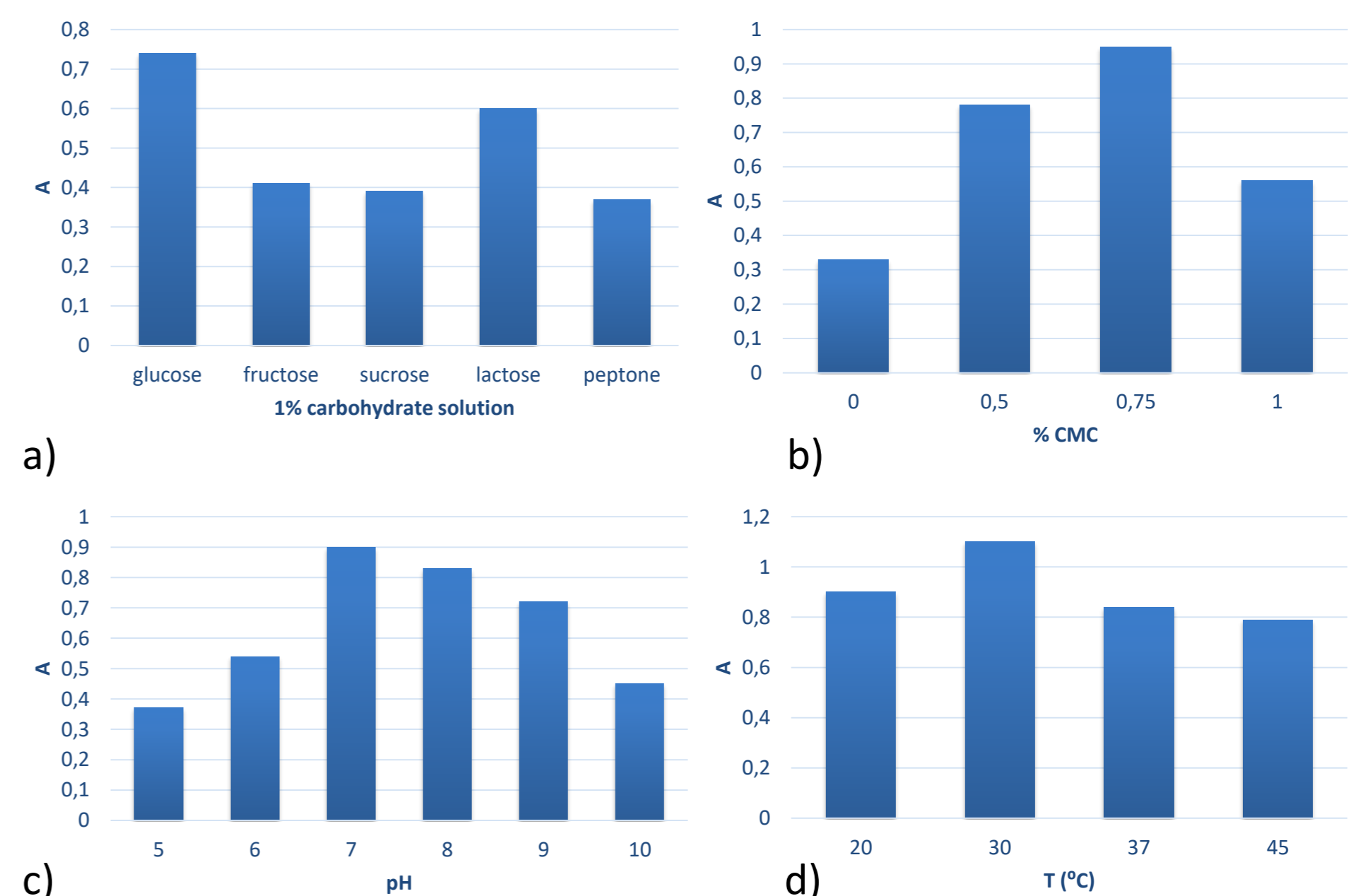


Figure 1. Optimization of the growing medium:

- a) percentage of carbohydrate source;  
b) percentage of carboxymethylcellulose (CMC); c) pH; d) temperature

Enzyme stability results showed that the enzyme retained 90% of its activity after minimal preincubation at pH 5.5 and more than 60% of its maximum activity at pH 4–10, demonstrating stability over a wide pH range. Therefore, it can be considered as an acid/alkali-resistant enzyme. It was found that the greatest activity of the enzyme occurs at 50 °C. When the temperature is increased to 50 °C, the enzyme activity increases and then decreases linearly. More than 60% of  $\beta$ -1,4-glucanase's enzymatic activity is observed in the range of 20-60 °C.

### CONCLUSION

Consequently, biocompositions based on *Lysobacter sp.* can be used in a wide range of temperatures and pH, which represents its promising use as a probiotic drug.

The work was carried out within the framework of the state task FZWG-2023-0007. Adaptive reactions of microorganisms: theoretical and applied aspects.

### FUTURE WORK / REFERENCES

In the future, it is planned to obtain biocompositions based on *Lysobacter sp.* and other probiotic microorganisms for the purpose of creating veterinary drugs and feed additives for farm animals.

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