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Growth optimization of *Pseudomonas L521* for bioremediation of caffeine contaminated environment

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Introduction

- Caffeine as a white crystalline alkaloid of xanthine that is bitter, odourless and amorphous in its pure form. At high concentration caffeine is toxic to saprophytic organisms that are concerned in important biotransformation in the environment, which causes disorder in environmental stability (Gibson et al., 2009; Ibrahim et al., 2015b)
- This growth optimization focuses on obtaining suitable parameters at which the caffeine-degrading bacterium can best function in degrading caffeine so that it can be used in bioremediation of caffeine contaminated environments without the use of chemicals.

Objectives

- To study the growth of *Pseudomonas sp.* in the presence of various carbon and nitrogen sources.
- To study the growth and biodegradability of Pseudomonas sp. at varied pH and temperature.
- To study the effect of different caffeine concentrations on the growth of *Pseudomonas sp*.

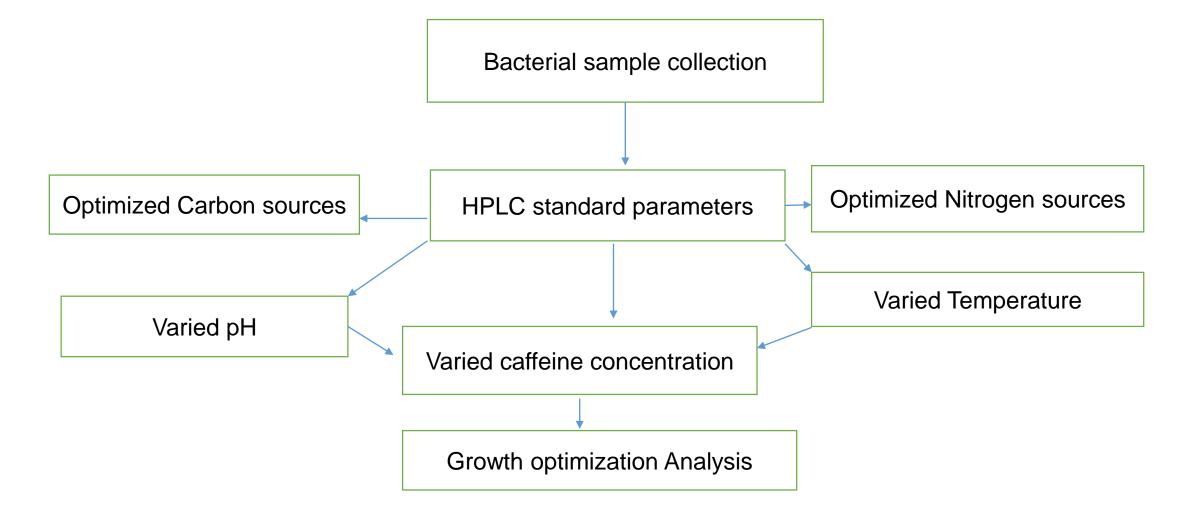


Methodology

- Sample collection and maintenance; Pseudomonas strain L521 was collected form Biological Science Department Nigerian Defence Academy, Kaduna and cultivated in a Caffeine Liquid Media (CLM). The bacterial strain was incubated on a rotary shaker incubator at 150rpm and 28 °C for 48 hrs to obtain a starter culture. The isolates were maintained and subcultured every 2 weeks in the caffeine agar medium.
- Growth Analysis; Caffeine degradation for the isolate was estimated by HPLC equipment.
- Growth optimization; Several parameters such as different concentration of caffeine, carbon source, nitrogen source, pH, and temperature were optimised in order to make sure that the caffeine degradation was successfully carried out for

BSP seudomonas strain

Experimental **Setting**





Results and Discussion

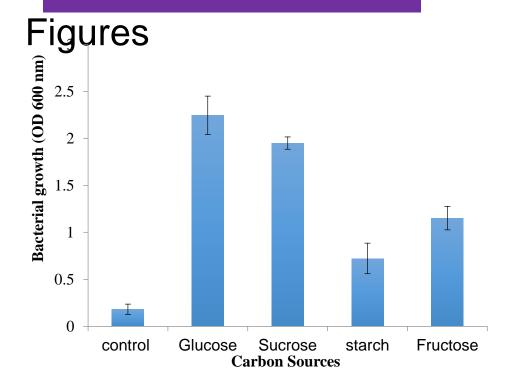


Figure 1: The effect of different carbon sources on bacterial growth and caffeine degradation of *Pseudomonas sp.* with initial caffeine concentration of 0.3 g/L, 5 g/L of carbon source and 0.4 g/L of ammonium chloride.

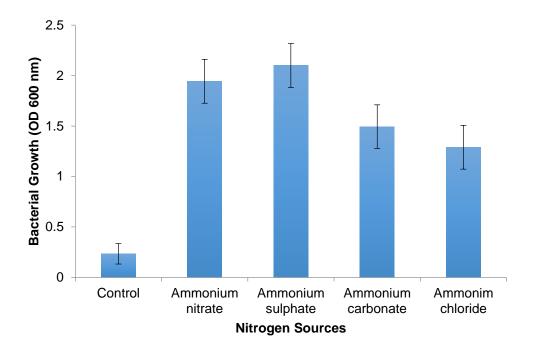


Figure 2: The effect of different nitrogen sources on bacterial growth and caffeine degradation of *Pseudomonas sp.* with initial caffeine concentration of 0.3 g/L, 0.4 g/L of nitrogen source and 5 g/L of sucrose.



Results and Discussion

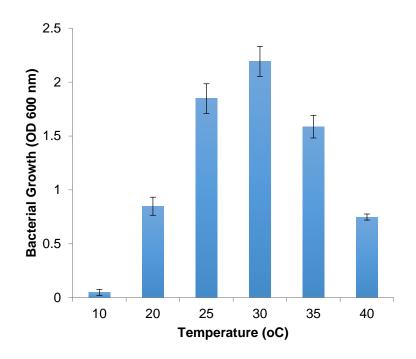


Figure 3: Effect of various temperatures on bacterial growth by *Pseudomonas sp.* with initial caffeine concentration of 0.3 g/L, 5 g/L of sucrose and 0.4 g/L of ammonium sulphate.

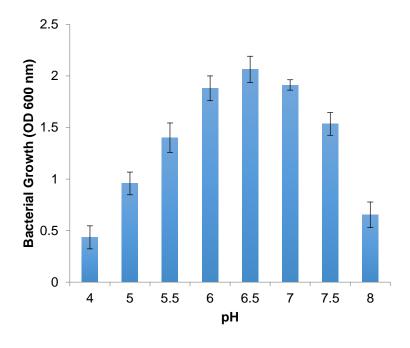


Figure 4: Effect of various pH on bacterial growth by *Pseudomonas sp.* with initial caffeine concentration of 0.3 g/L, 5 g/L of sucrose and 0.4 g/L of ammonium chloride.

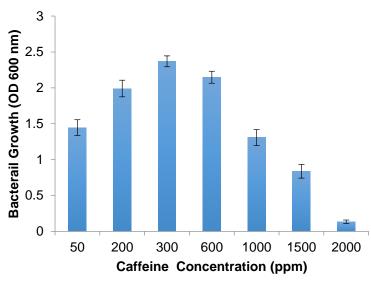


Figure 5: The effect of different caffeine concentrations on bacterial growth and caffeine degradation of *Pseudomonas sp.* with initial caffeine concentration of 0.5 mg/L, 5 g/L of carbon source and 0.4 g/L of ammonium chloride.



Conclusions & Recommendations CONCLUSION

• The aim of this research was to evaluate biodegradability of Pseudomonas L521 on caffeine. The growth optimization studies provides the optimum growth conditions for Pseudomonas sp. that can be used to increase the capability of the bacteria in degrading caffeine. Pseudomonas sp. grew best at 5g/L glucose, 0.4g/L ammonium sulphate, pH 6.5, 30 °C and 0.3g/L caffeine.

RECOMMENDATION

- Pseudomonas sp was found to be the best caffeine degrading bacteria from the comparison of the literature and the result in this research.
- Therefore, further studies on cloning the gene, purification and hyper-expression of enzyme and its molecular techniques would be essential.

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