

*Conference Name: BioTecnica 2026 – International Conference on Advances in Biological Sciences,*

*21-22 January, Ho Chi Minh*

*Conference Dates: 21-Jan- 2026 to 22-Jan- 2026*

*Conference Venue: PARKROYAL Saigon, 311 Đ. Nguyễn Văn Trỗi, Phường 1, Tân Bình, Hồ Chí Minh, Vietnam*

*Appears in: LIFE: International Journal of Health and Life-Sciences (ISSN 2454-5872)*

*Publication year: 2026*

*Sunday Stephen Abi, 2026*

*Volume 2026, pp. 01-02*

*DOI- <https://doi.org/10.20319/icrlsh.2026.0102>*

*This paper can be cited as: Abi, S. S.(2026). Temperature-Dependent Antimicrobial Elicitation in Environmental Bacteria Co-Cultured With *Enterococcus faecalis* in the Presence of Ampicillin/Tetracycline. BioTecnica 2026 – International Conference on Advances in Biological Sciences, 21-22 January, Ho Chi Minh. Proceedings of Healthcare and Biological Sciences Research Association (HBSRA), 2026, 01-02*

## **TEMPERATURE-DEPENDENT ANTIMICROBIAL ELICITATION IN ENVIRONMENTAL BACTERIA CO-CULTURED WITH *ENTEROCOCCUS FAECALIS* IN THE PRESENCE OF AMPICILLIN/TETRACYCLINE**

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### **Abstract**

**Research Objectives:** *This study investigated whether co-culturing environmental bacterial isolates against *Enterococcus faecalis* (NTUCC 687) in the presence of ampicillin or tetracycline could elicit antimicrobial production, and whether this response was temperature-dependent.*

**Methodology:** *A total of 96 environmental bacterial isolates (arrayed in a 96-well plate) collected from built and natural surfaces around NTU Clifton campus were co-cultured against an *Enterococcus faecalis* lawn using an in-house stamping assay on UTI ChromoSelect agar. Plates were incubated at 25°C, 30°C, 37°C, and 42°C for 24–72 hours. For elicitation, UTI agar was fortified separately with ampicillin (0.05 µg/mL) or tetracycline (0.05 µg/mL). These low (sub-inhibitory) concentrations were used as*

*stressors to probe signalling-associated induction of antimicrobial activity rather than direct growth inhibition. Antimicrobial activity was recorded as categorical zones of inhibition, classified as small (+), medium (++) or large (+++).*

**Findings:** *Tetracycline elicited a higher frequency of large (++) zones of inhibition than ampicillin, peaking at 30°C (12 isolates, 12.5%), followed by 42°C (9 isolates, 9.4%), 25°C (5 isolates, 5.2%), and 37°C (3 isolates, 3.1%). In contrast, ampicillin produced only two large zones (2.1%), both observed at 25°C. These patterns indicate that antimicrobial elicitation was modulated by both antibiotic stressor type and incubation temperature within the co-culture assay.*

**Research Outcomes and Future Scope:** *These findings support co-culture-based elicitation as a practical strategy for uncovering latent antimicrobial potential in environmental bacteria and highlight temperature as a key modulator of expression. Future work should identify the active producers and compounds involved, investigate underlying molecular mechanisms, and assess scalability for antimicrobial discovery.*

**Keywords:**

Antimicrobial Elicitation, Co-Culture, *Enterococcus faecalis*, Antibiotic Stress, Temperature