14th International Conference on Plant Pathogenic Bacteria (ICPPB)

ABSTRACT SUBMISSION FORM

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[] Disease Epidemiology and Pathogen Ecology
[] Disease Emergence and Pathogen Evolution
[] Disease Control and Prevention
[] New Tools in Disease Diagnostics and Pathogen Identification
[1] Molecular Plant - Bacteria (and Insect) Interactions
[] Bacterial Pathogens and the Phytobiome
[] Natural and Engineered Plant Disease Resistance

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The *iaaL* gene in the *Pseudomonas syringae* complex: functional characterization and biological activity

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Phytopathogenic bacteria of the *Pseudomonas syringae* complex are causal agents of diseases in a wide variety of woody and herbaceous plants with agronomic and ornamental interest. Indole-3-acetic acid (IAA) is an auxin phytohormone whose production is widely distributed among plant-associated bacteria. Some P. syringae strains can further metabolize IAA to the amino acid conjugate 3-indole-acetyl-ɛ-L-lysine (IAA-Lys), a process involving the enzyme IAA-Lys synthase, encoded by the *iaaL* gene. IAA-Lys is less biologically active than IAA, so it has been speculated that the conjugation of IAA with L-Lysine could allow the bacteria to control the levels of free IAA accumulated in the bacterial cytoplasm and/or secreted to the plant tissues. The *iaaL* gene is widespread in the *P. syringae* complex, and three different alleles (iaaLPsv, iaaLPsn and iaaLPto) have been described [1]. Recently, we have identified a fourth allele (iaaLPsf) specifically encoded in the genome of strains isolated from Fraxinus excelsior. However, comparative analyses of the biochemical and biological activities of the different iaaL alleles have not been performed. In this work, the genomic context of these four alleles in a collection of *P. syringae* complex strains has been analyzed. In addition, we have constructed strains overexpressing each of these *iaaL* alleles and analysed their biological activities using an elongation assay of Arabidopsis thaliana roots. Finally, expression of these alleles in E. coli allowed the purification of these four IaaL proteins and the analysis of their specific activities using an *in vitro* enzymatic coupling assay.

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[1] Matas et al., 2009. Appl. Environ. Microbol., 75, 1030-1035.

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