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New insights into indole-3-acetic acid metabolism in *Azospirillum brasilense*.

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Abstract

AIMS: The aim of this research was to analyze the global IAA metabolism in three commercially used strains of *A. brasilense*.

METHODS AND RESULTS: *A. brasilense* Sp245, Az39 and Cd, containing a plasmid with the ipdC-gusA fusion (pFAJ64), were cultured in minimal medium MMAB with or without 10 mg.l⁻¹ of L-trp till exponential or stationary growth phase. The cultures were then split into 10 ml tubes and individually treated with 10 mg.ml⁻¹ IAA, IBA or NAA (auxin catabolism and homeostasis); IAPhe, IALeu, IAA-ala, IAA-glucose (IAA conjugate hydrolysis); or L-lys, L-leu, L-ileu, L-phe, L-ala, L-val, L-arg, L-glu, L-his, L-met, L-asp, L-cys, L-ser, L-pro, L-thr and L-trp (regulation of IAA biosynthesis and IAA conjugation). Bacterial growth, IAA production and ipdC expression were evaluated. None of the *A. brasilense* strains were able to hydrolyze IAA conjugates, catabolize auxins, or conjugate IAA with amino acids or glucose. L-amino acids L-met, L-val, L-cys and L-ser inhibited bacterial growth and decreased IAA biosynthesis. The expression of ipdC and IAA biosynthesis but not bacterial growth were affected by L-leu, L-phe, L-ala, L-ile, L-pro. L-arg, L-glu, L-his, L-lys, L-asp and L-thr did not affect any of the measured parameters.

CONCLUSIONS: In this paper we confirmed that *A. brasilense* produces IAA only in presence of L-trp, is not able to degrade auxins, conjugate IAA with sugars and/or L-amino acids, or hydrolyze such conjugates to release free IAA. Finally, we found that bacterial growth and/or IAA biosynthesis were inhibited by the presence of several L-amino acids probably by diversion of the cellular metabolism.

SIGNIFICANCE AND IMPACT OF STUDY: We propose a renewed model to explain IAA metabolism in *A. brasilense*, one of the most studied phytostimulatory bacteria. This article is protected by copyright. All rights reserved.

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KEYWORDS: *Azospirillum brasilense* ; L-amino acids; auxins; catabolism; conjugation; homeostasis; hydrolysis; indole-3-acetic acid; metabolism

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