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EXPRESSION OF A CHIMERIC ANTIMICROBIAL PROTEIN IN TRANSGENIC TOMATO CONFERS RESISTANCE TO THE PHYTOPATHOGEN *Ralstonia solanacearum*¹ / A expressão de uma proteína quimérica antimicrobiana em tomate confere resistência às plantas contra *Ralstonia solanacearum*. <u>T.P. MORAIS</u>²; R. NASCIMENTO²; L.R. GOULART²; J.M.Q. LUZ²; A.M. DANDEKAR³. ²Universidade Federal de Uberlândia, Uberlândia, Brasil; ³University of California, Davis, USA. E-mail: morais_prado@hotmail.com

Research interest on antimicrobial peptides has increased because of their broad range activity, resulting in several biotechnological applications addressed to plant protection. The present study taps into the *in vitro* characterization of a chimeric protein and its potential use for development of transgenic tomato plants with resistance to a bacterial pathogen. The chimera was designed based on the NE-CecB antimicrobial protein, which has been previously validated on the plant pathogen Xylella fastidiosa. Each domain was substituted by homologous genes found in plant genomes, comprising a pathogenesis-related protein (SIP14a) joined to a plant-derived cecropin B-like peptide (an α -helix from phosphoenolpyruvate carboxylase - PPC20). In vitro antibacterial activities of SIP14a and SIP14a-PPC20 were confirmed in kill-curves assays against the bacterial wilt pathogen Ralstonia solanacearum, suggesting their use as promising candidates in plant protection. Later, tomato plants were engineered to express SIP14a-PPC20 chimera and challenged with R. solanacearum. Within control plants (wild-type Solanum lycopersicum cv. Money Maker), disease evolved from wilting symptoms to plant death in two weeks. SIP14a-PPC20-transgenic plants, however, showed no symptoms or reduced disease severity. Bacterial multiplication in stems of transgenic plants was suppressed more than 2-fold compared to control plants, and absence of disease symptoms could be associated with this growth suppression.

Keywords: Genetic engineering; Disease resistant plants; Therapeutic antimicrobial proteins.

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