

## Overexpression of the *lb-rolB/C* gene perturbs biosynthesis of caffeoylquinic acid derivatives in transgenic calli of sweet potato

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*Ipomoea batatas* is a root crop widely cultivated in South America, Africa, and Asia. This is also a source of caffeoylquinic acid derivatives (CQAs) with potential health-promoting benefits. Sweet potato genome carries two separate cellular T-DNA (cT-DNA) regions (*lbT-DNA1* and *lbT-DNA2*). Especially, *lbT-DNA2* contains five ORFs homologous to the *Agrobacterium rhizogenes* T-DNA, namely ORF13, ORF14, ORF17n, ORF18/ORF17n, and RolB/RolC proteins [1]. Unfortunately, presently there is insufficient information on *lbT-DNA2* genes' function in the physiological processes of sweet potatoes.

In this study, expressional levels of the *lbT-DNA2* genes and the effect of *lb-rolB/C* overexpression were examined using *I. batatas* cell culture. We discovered that *I. batatas* cT-DNA genes were not expressed in callus, and abiotic stresses and chemical elicitors affected their transcriptional levels weakly. Additionally, two *lb-rolBC*-transgenic cell lines have been established though *Agrobacterium*-mediated transformation of *I. batatas* callus cells. Overexpression of *lb-rolB/C* gene reduced biomass accumulation of transgenic cell lines by 1.2–1.6 times and increased the CQAs content by 1.5–1.9-fold. To justify the metabolic fluctuations, the study also looked into the expression patterns of the major biosynthetic genes, namely *bPAL*, *lbC4H*, *lb4CL*, *lbHCT*, and *lbHQT*. The obtained data demonstrated that the overexpression of the *lb-rolB/C* reduced the *lbPAL* transcript but considerable increase in the transcript levels of the *lbHQT*. We propose that this result was obtained through as-yet-uncharacterized signaling pathways activated by RolB/RolC.

### REFERENCE

1. Kyndt T, Quispe D, Zhai H, et al. The genome of cultivated sweet potato contains *Agrobacterium* T-DNAs with expressed genes: an example of a naturally transgenic food crop. *PNAS USA*. 2015;112(18):5844–5849. DOI: 10.1073/pnas.1419685112

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