

**SCREENING OF SOLANUM LYCOPERSICUM X SOLANUM PENNELLII
INTROGRESSION LINES FOR THE RESISTANCE VERSUS THE PARASITIC WEED
PHELIPANCHE RAMOSA: A STEP TOWARDS DUAL RNA-SEQUENCING AIMED AT
STUDYING HOST/PARASITE CROSSTALK**

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Broomrapes (*Orobanche* spp. and *Phelipanche* spp.) cause severe damages to crops every year around the world. Particularly, *Phelipanche ramosa* is a threat that severely affects tomato (*Solanum lycopersicum* L.) cultivation. One of the most effective approaches to control this pest is to develop resistant varieties through breeding. Unfortunately, within the cultivated tomato gene pool there are scarce sources of resistance, thus significantly impairing resistance versus this natural enemy. However, some preliminary and recent evidence indicates that *Solanum pennellii*, a close wild relative of *S. lycopersicum*, is resistant. Some *S. lycopersicum* x *S. pennellii* introgression lines (ILs) among those available may show considerable tolerance to parasitic weed infestations.

Interestingly, preliminary results on root exudates of *S. pennellii* showed the absence of the classical strigolactones usually found in cultivated tomatoes (i.e., orobanchol, solanacol and didehydro-orobanchol).

The screening for the resistance of ILs versus *P. ramosa* is ongoing. We are testing the ability of tomato plantlets to stimulate *P. ramosa* seed germination and induce the production of the haustoria and the development of the tubercles, by performing in vitro germination assays and plastic bag assays. The purpose of these experiments is to identify the most resistant and most susceptible ILs to be subjected to dual RNA-Sequencing. Indeed, host/parasite interaction determines large-scale reprogramming of transcriptomes in both organisms. The analysis of transcriptome dynamics and gene expression profiles is essential to reveal the molecular mechanisms that mediate host/parasite crosstalk and to eventually identify the key genes involved in the defense strategies implemented by the host.

In addition, some lines of evidence suggest that non-coding RNAs (ncRNAs) play an active role in the interaction between tomato and different pathogens and that in other species long ncRNAs are transferred bidirectionally through vascular connections acting as critical regulators in coordinating host/parasite interaction. Tomato root samples (3 biological replicates) and samples of *P. ramosa* at different stages of infection (pre-attachment, haustorium differentiation, tubercle development) will be collected, the total RNA extracted and then sequenced to simultaneously study both the evolution of the parasitic weed and the host response. The analysis of the coding and non-coding component (never investigated before) of the transcriptome could lead to expanding the panel of candidate genes useful as targets in future mutagenesis programs to obtain tomato varieties resistant to Orobanchaceae.