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Two-hybrid assay in plants

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In the August issue of *Nature Biotechnology*, Rajagopal Subramaniam and colleagues from the *Université de Montreal* describe a system for visualizing protein-protein interactions in living plant cells (*Nature Biotechnology* 2001, **19**:769-772). Their technique is based on an *in vivo* **protein-fragment complementation assay** in which protein interactions between fusion proteins induce folding and reassembly of fragments of the murine dihydrofolate reductase (DHFR) enzyme. Protein-protein interactions are monitored by the DHFR inhibitor fluorescein-conjugated methotrexate (fMTX). Subramaniam *et al.* used their assay to examine the interaction between *Arabidopsis* disease-resistance protein **NPR1/NIM1** and the basic leucine-zipper protein TGA2. They show that salicylic acid induced NPR1-TGA2 interaction in tobacco or potato cells. This plant two-hybrid system should prove useful for the functional annotation of plant genomes.

References

1. *Nature Biotechnology*, [<http://www.nature.com/nbt/>]
2. Université de Montreal , [<http://www.umontreal.ca>]
3. Clonal selection and *in vivo* quantitation of protein interactions with protein-fragment complementation assays.
4. The *Arabidopsis* *NPR1* gene that controls systemic acquired resistance encodes a novel protein containing ankyrin repeats