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Different methods of evaluation of Monilinia laxa on apricot flowers and branches

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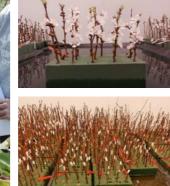
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State of the art

- ✓ Organic apricot production is currently not profitable.
- ✓ The main obstacle to sustainable profitability is brown rot caused by the fungus Monilinia Iaxa (Aderh. & Ruhl).
- ✓ In the current apricot germplasm no source of total resistance has been shown, but some varieties are expressing interesting levels of tolerance.
- ✓ A good evaluation of the M. laxa symptoms is essential for a precise diagnosis of the infection and to appreciate differences between tolerant and susceptible varieties and genotypes





Results and Conclusions

- ✓ Different levels of infection were observed within the bi-parental population for the three controlled phenotyping evaluations.
- ✓ Symptoms in petals were not linked with the Monilinia infection according to our observations. The infection with the spores on the pistil test was very low.
- ✓ Good segregation was observed for the visual assessment and for the evaluation test on branches
- ✓ There was not a good correlation between the visual assessment in the field compare with the evaluation of branches and flowers.
- \checkmark There was not a good correlation neither between the three evaluations under controlled conditions.
- ✓ Based onto the observed variability, a QTL approach can be applied for assessing the genetic components involved in Monilinia resistance.



Materials and methods

- ✓ Different evaluation methods were carried out on a bi-parental population between Bakour (tolerant to M. laxa) and Bergeron (susceptible):
- 1) Visual evaluation of Monilinia symptoms was carried on the trees (from 0 to 100% of infection) 35 days after full blossom. Wheater dependant method.
- 2) Evaluations Under controlled conditions on each genotype:
 - a) A spore suspension (104 / ml) of *M. laxa* was sprayed on flowers (20°C, 90% HR, 14 hours day), and % of infected flowers (necrotic petals) was measured 36 hours after.
 - b) A drop of spores (10⁵/ml) was inoculated with a pipette directly on the pistil when the flower was in stage F. Flower / branches infection was evaluated.
 - c) A plug of M. laxa mycelium was added on branches (20°C, 80% HR, darkness), and 8 days after the length of the reaction was measured.

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