

Differences between the Xanthomonas arboricola pv. pruni genome and the model strains Xcc B100 and Xcv 85-10 using DNA microarray technology

Autor(es):	MAYER, Lauri; BECKER, Anke; PÜHLER, Alfred; SILVA, Wladimir Padilha;
	VENDRUSCOLO, Claire Tondo
Apresentador:	Lauri Mayer
Orientador:	Wladimir Padilha da Silva
Revisor 1:	Marcelo Mendonça
Revisor 2:	Caroline Bastos
Instituição:	Universidade Federal de Pelotas

Resumo:

The phytopathogenic bacteria Xanthomonas arboricola pv. pruni is the causal agent of Prunus Bacterial Spot disease (PBS) that infects cultivated Prunus species and their hybrids. In southern Brazil this pathovar is an important pathogen in peach and plum crops, causing considerable economic losses annually. On the other hand, most of the Xanthomonas species including X. arboricola pv. pruni produces by an aerobic fermentation process the xanthan gum, an important biopolymer used in the food, oil and cosmetic industry. To the Comparative genomic Hybridisation five strains isolated from peach and plum trees at the Centro de Pesquisas Agropecuárias de Clima Temperado (EMBRAPA, Pelotas, RS, Brazil) were used. For this study were choice strains with different capacity for xanthan production (data not shown) in MPII medium. The genomic DNA was hybridized with the model strains X. campestris pv. campestris B100 and X. campestris pv. vesicatoria 85-10 using the microarray technology. Data analysis was done by applying the ImaGene 6.0 software (Biodiscovery Inc., Los Angeles, CA) for acquisition of the mean signal for each spot of the microarray and the EMMA 2.2 software (Bielefeld University, Germany) for normalization and t-statistics. A gene was considered to have a statistically significant difference if the P value was ≤ 0.01, the M value (log2-ratio) was ≥ 2 or ≤ -2 and the A value (mean intensity) was ≥ 7. The results shown a high genetic similarity among the five strains used. Several gene clusters present in the X. campestris pv. campestris B100 and X. campestris pv. vesicatoria 85-10 genome were found to be missing in the X. arboricola pv. pruni genome. Of these, were identified VirB genes (VirB1, virB2, virB5, virB8, virB9, virB11), icm genes (icmB, icmC, icmJ, icmT) and pil genes (pilC, pilE, pilL, pilO, pilQ, pilR, pilX, pilY, pilV). VirB and icm genes belong to the type IV secretion system, which is used to transport toxic molecules to host cells and also to DNA or protein-DNA complexes transport. The genes pilX, pilY and pilV are Tfp pilus assembly proteins and the other pil genes are involved with secretion systems and membrane proteins. These pilus assembly proteins are important for virulence and DNA transfer in a variety of prokaryotes. In conclusion, the absent of the VirB, icm and pil gene groups in the X. arboricola pv. pruni genome are probably partially responsible of the low pathogenicity of this pathovar.