

## ***Molecular properties of Cherry leaf roll virus***

J. Langer, S. von Bargen, C. Büttner

Faculty of Agriculture and Horticulture, Section Phytomedicine, Humboldt-Universität zu Berlin, Lentzeallee 55/57, D-14195 Berlin, Germany

Contact: langerj@rz.hu-berlin.de

The *Cherry leaf roll virus* (CLRV) is a globally distributed pathogen occurring primarily on deciduous, fruit and ornamental trees from at least 17 genera, including many economically important trees like walnut, cherry and birches. CLRV is a nepovirus of the *Comoviridae* within the *Picornavirus* superfamily with a bipartite genome organisation and protein expression strategy resembling other members of the genus. Nepoviral RNAs exhibit 3' non-coding regions (3' NCR) with extensive sequence identities (80-100 %), exclusively illustrated by the members of the nepovirus subgroup c, including the CLRV, with very large 3' NCRs of over 1500 nt. Sequence comparisons between the RNA1 and RNA2 specific 3' NCRs of six different CLRV isolates from different host plant species and phylogenetic groups displayed almost identical 3' NCRs (97.5-99.5 %) for five CLRV isolates. A raspberry isolate exhibits 3' NCRs with only 73.8 % sequence identity, raising the question about the prerequisite of sequence identity within the 3' NCRs of a RNA population of an individual CLRV strain. So far, the question for the benefit of the long 3' NCRs in any replication or translation mechanism is still unanswered, but the selective 3' NCR sequence conservation of almost all previously analyzed nepovirus isolates, confirmed a strict necessity of identity for maintaining functional sequences within this region. It is commonly considered that homologous recombination is responsible for the 3' terminal sequence identity. But this is only one of several efficient mechanisms to ensure viability of RNA populations, at least for the CLRV since a raspberry isolate with non-homologous 3' NCRs was found in this study. Furthermore, a stable secondary hairpin structure was predicted within the analyzed 3' NCRs of all six different CLRV isolates. This is located in a region with high sequence variability of up to 34 % and the conservation of this secondary structure suggests that it represents an important functional domain within the 3' terminus of CLRV-RNAs.