

Analysis of the resistance breaking ability of different beet necrotic yellow vein virus isolates loaded to a single *Polomyxa betae* population in soil

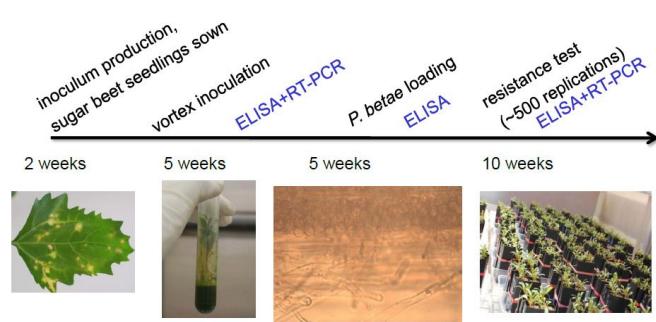
Introduction

Beet necrotic yellow vein virus (BNYVV) is vectored by *Polomyxa betae* and causes rhizomania in sugar beet. By cultivating partial resistant genotypes the disease can be controlled. Resistance genes like *Rz1* and *Rz2* reduce virus replication and prevent virus spread from infected hair roots into the tap root. The RNA 3 encoded pathogenicity factor P25 is responsible for symptom expression and virus translocation in the root system. P25 has a highly variable amino acid composition at amino acid tetrad 67-70. The uniformly applied *Rz1* resistance is suggested to exert a positive selection on the P25 tetrad composition because isolates with the ability to overcome *Rz1* display a valine at amino acid position 67 (V67) (Koenig et al., 2009). However, direct comparison of aggressive isolates has remained impossible due to varying vector population densities and other soil-borne pathogens.

Material and Methods

A method to load a virus-free *P. betae* population was developed (Fig. 1). Three different BNYVV isolates with different tetrad compositions (A-ACHG, IV-VCHG, P-SYHG) were used. Vortex inoculation method was adapted and modified after Koenig and Stein (1990).

Fig. 1: Material and methods used to load a *P. betae* population



Results

Tab. 1: Number of infected plants (%) after vortex inoculation and resistance test

BNYVV isolate	Genotype	Vortex inoculation	Resistance test
		% infected plants ^a	% infected plants ^b
A	<i>rz1rz1</i>	96	100
A	<i>Rz1rz1</i>	na ^c	45
A	<i>Rz1rz1+Rz2rz2</i>	na	4.5
IV	<i>rz1rz1</i>	96	100
IV	<i>Rz1rz1</i>	na	87.5
IV	<i>Rz1rz1+Rz2rz2</i>	na	2.2
P	<i>rz1rz1</i>	84	92.1
P	<i>Rz1rz1</i>	na	73.7
P	<i>Rz1rz1+Rz2rz2</i>	na	7.5

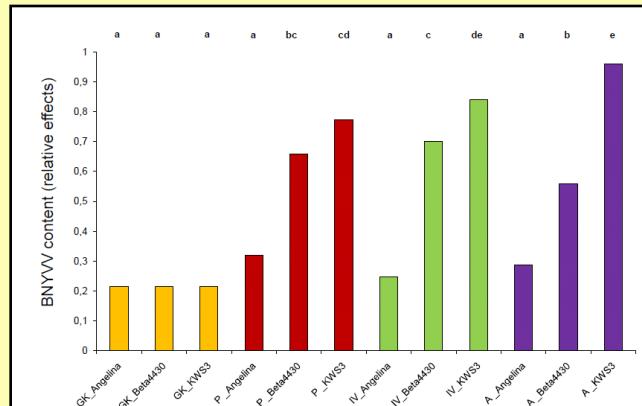
^a mixed sample obtained from seven plants per pot; ^b mixed sample obtained from four plants per pot; ^c na = not applicable

- A virus-free *Polomyxa betae* population from a naturally infested field soil was loaded.

- A high infection efficiency was achieved (Tab. 1).

- In order to evaluate the data properly, nonparametric analyses was performed and results shown as relative effects (Fig. 2).

Fig. 2: Total virus contents induced by three BNYVV isolates (A, IV, P) on different sugar beet genotypes shown as relative effects



References: Koenig, R. and Stein, B. (1990). Schriftenreihe der Deutschen Phytomedizinischen Gesellschaft, vol. 1. Proceedings of the First Symposium of the International Working Group on Plant Viruses with Fungal Vectors, Braunschweig, Germany, August 21-24, pp. 87-90.
Koenig, R., Loss, S., Specht, J., Varrelmann, M., Lüddecke, P., and Deml, G. (2009). J. Gen. Virol. 90:759-763.