**Evaluating Low Levels of Tachigaren on Minimum Build-Up-Treated Sugar Beet** Seed for Protection Against *Aphanomyces cochlioides*. R. M. Harveson, L. Hubbell, C. E. Windels, J. A. Smith, J. R. Brantner, J. F. Giles, and N. R. Cattanach.

The objective of this study was to evaluate low levels of Tachigaren (hymexazol) as possible standard treatments on sugar beets under low to moderate levels of disease pressure, caused by *A. cochlioides*. The study was conducted for 3 years (2001-2003), at 12 separate sites in Michigan, Nebraska, and North Dakota. Treatments consisted of 1) Apron /Thiram incorporated into minimum build-up coatings with Tachigaren (20g and 30g), 2) Tachigaren applied at 45 g in a standard seed pellet, and 3) Apron/Thiram-treated controls. The same treatments were additionally tested under optimum conditions in the greenhouse in field soils naturally infested with varying concentrations of *A. cochlioides*. Field results varied, but several locations showed higher rates of Tachigaren with minimum build-up treatments (30g) caused reduced seedling emergence. However, few significant differences were observed from yield parameters, suggesting minimal damage to crop at the end of the season. Greenhouse results indicated that using low rates of Tachigaren with minimum build-up coatings exhibited more potential for use in fields with low-moderate levels of A. cochlioides. These same treatments were not consistent in soils with high disease potential.

This same mutation has been shown to confer sensitivity to N-phenyleubamates (NPC). When our C besicola isolates were tested for sensitivity to the NPC tongocide diethofencare (DPC), benzimulazole-tolerant isolates proved sensitive to the NPC, while bacainidazole-sensitive isolates showed little or no a cusitivity to DPC. Ninety-six C 2004. These isolates were collected in the central High Plains region of the United States in besicola isolates were collected in the central High Plains region of the United States in 2004. These isolates were tested for their response to benzimidazoles and DFC and forty-three (45%) mare highly sensitive to MB (100% growth inhibition at 1 ppm) and befenant to DFC. Fifty-three (55%) were highly interast to MB and did not growth of 0.971. PCR primers for the benzimidazole-tolerani and benzimidazole-sensitive sequences are being developed to investigate use in a more tipid method for detection of benzimidazole sensitivity. The negative cross resistance and hnowledge of the mutation benzimidazole sensitivity. The negative cross resistance and hnowledge of the detection of benzimidazole sensitivity. The negative cross resistance and hnowledge of the mutation benzimidazole sensitivity. The negative cross resistance and knowledge of the detection of benzimidazole sensitivity. The negative cross resistance and knowledge of the mutation benzimidazole sensitivity.





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