PROGRESS REPORT TO THE MAINE POTATO BOARD January 26, 2015

Project Title

Building Varietal Resistance to Late Blight and PVY using Marker-assisted Selection

Investigators

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Executive Summary

Improved variety resistance will be a key long-term means of managing the new strains of both late blight and PVY that are plaguing the industry. Marker-assisted selection is an effective method of speeding the development of new potato varieties with key disease resistance traits; however, only a few useful molecular markers are currently available to help breeders select potato clones that have desirable traits. In this project, we are developing late blight and PVY resistant potato clones through the use of marker-assisted selection. We will utilize phenotyped and genotyped late blight resistant parental clones to begin identifying likely resistance genes. We will begin developing genetic markers that can be used to speed future breeding, selection and variety development for late blight resistance. When the resulting materials also have a PVY resistant parent in their background, we will use two PVY resistance markers (RYSC3 for Ry_{adg} ; YES3 for Ry_{sto}) to speed identification of new PVY-resistant potato clones. Where appropriate we will also use the H1 marker for golden nematode resistance. Maine Potato Board support during 2012 and 2013 allowed us to successfully incorporate the RYSC3 and H1 markers into our breeding efforts. The marker-assisted approach will help us identify resistant clones earlier in the program, thus increasing the chances of retaining resistant clones over several years of field selection and improving the program's chances of releasing useful varieties with resistance to late blight, PVY, and golden nematode. This research will also allow us to more quickly identify resistant clones for use in future crossing programs designed to generate resistant varieties.

Duration of Project

Funding was requested for April 1, 2014 to March 31, 2015 to support the expansion of marker-assisted disease resistance breeding as an integral component of the potato breeding pipeline at Aroostook Research Farm. The goal is to develop simple and effective DNA-based procedures for genotyping previously established breeding lines, new clone selections, and their derivatives for late blight resistance.

Project Objectives:

To reduce the impact of late blight through the development of new varieties and breeding lines with effective resistance to diverse populations of the late blight pathogen

To use DNA marker-assisted selection strategies for stacking known genes for late blight and potato virus-Y resistance (e.g., RYSC3 for Ry_{adg} , YES3 for Ry_{sto}) in common genetic backgrounds that are economically important to the potato industry in Maine and the eastern United States

Grant Received:

\$6,000 (\$12,000 was requested)

Progress on late blight marker-assisted selection

Late blight resistance is one of the priorities of The University of Maine Potato Breeding Program. Heretofore, resistant clones and varieties have been identified by phenotypic selection against field populations of late blight. With this research, we are moving toward a more strategic, genetic-based selection process that maximizes opportunities for 'pyramiding' resistance genes. Resistant clones and varieties from within the ME program and from the USDA-ARS Beltsville potato breeding program have been identified as important candidates. Example late blight resistance sources that are being examined include: B0718-3, Barbara, Dakota Trailblazer, Defender, Dorita, Elba, J117, Jacqueline Lee, NY121, Missaukee, Pirola, Stirling, Tollocan, Torridon, and Yukon Gem. In all, more than 60 varieties and clones will be studied.

Protocols for eleven resistance genes derived from *Solanum bulbocastanum*, *S. demissum*, *S. microdontum*, and *S. phureja* have been gathered from the literature. All available markers for *S. demissum R*-genes will be utilized: *R1*, *R2*, *R3a*, *R3b*, *R8*, and *R10*. The *R1* and *R2* genes have been completely sequenced, so the identification of these genes should be very reliable. The protocols for the other four genes are associated with nearby chromosomal markers. The protocols for *R3a*, *R3b*, and *R8*, were used in a study to show that the Mastenbroek differential clones MaR8 and MaR9 are actually stacked with multiple late blight genes derived from *S. demissum* (Kim et al., 2012). The marker associated with the *R10* gene is 0.05 cM away, indicating extreme likelihood that the gene is present when the marker is. The new *S. bulbocastanum* hybrid showing aphid resistance as well as late blight resistance, Sbu8.5, is known to have the *RB/Rpi-blb1* gene, and will be tested for the *Rpi-blb2* and *Rpi-blb3* genes. All three genes have been completely sequenced, so these markers will be highly reliable. The results of this study will be the start of a late blight genetic profile for each variety and clone.

DNA extractions will begin later this winter, with PCR-based marker testing immediately

following. Additionally, almost 40 of these varieties and clones will be phenotyped against specific late blight races (e.g. US-23, US-24, etc.) in greenhouse or growth chamber trials and leaf assays. The varieties and clones will be measured for disease amount and rate. These experiments will show specific race resistance in these popular breeding varieties and clones.

The results of the genetic and phenotypic data of each variety and clone will be combined to examine the relationships between resistance and certain gene combinations. While many combinations of resistance genes have been shown to be additive, a study from October of last year indicates the possibility that some resistance genes interact to suppress resistance (Stirnweis et al., 2014). The authors suggested that this was a reason that a resistance gradient exists in resistance crosses. If there are any patterns of suppressive and additive resistance, we will be able to detect them when we look at the genotyping and phenotyping data together. The outcomes of this analysis will highlight the effectiveness of certain gene combinations, and may help potato breeders be more efficient at the crossing block.

Progress on PVY marker-assisted selection

Resistance to PVY has become a high priority in The University of Maine Potato Breeding Program as concerns about the disease have increased in the industry. With the reduced funding that was available for the proposed project, we focused our efforts on new marker development for late blight resistance. Where resistance sources are also likely sources of PVY resistance, we will use DNA-based marker technology to evaluation the likelihood of stacked late blight and PVY resistance genes; however, we have moved the routine use of DNA-marker based technology for selection of PVY resistant clones to other funding sources and will report on that effort in our overall potato breeding program report.

References

- Kim, H-J., H-R. Lee, K-R. Jo, S.M. Mahdi Mortazavian, D.J. Huigan, B. Evenhuis, G. Kessel, R.G.F. Visser, E. Jacobson, and J.H. Vossen. 2012. Broad spectrum late blight resistance in potato differential set plants MaR8 and MaR9 is conferred by multiple stacked R genes. *Theoretical and Applied Genetics* 124: 923-935.
- Stirnweis, D., S.D. Milani, S. Brunner, G. Herren, G.Buchmann, D. Peditto, T. Jordan, and B. Keller. 2014. Suppression among alleles encoding nucleotied-binding-leucine-rich repeat resistance proteins interferes with resistance in F₁ hybrid and allele-pyramided wheat plants. *The Plant Journal* 79: 893-903.