

**USDA-ARS/  
U.S. Wheat and Barley Scab Initiative  
FY11 Preliminary Final Performance Report  
July 13, 2012**

**Cover Page**

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<b>Fiscal Year:</b>	FY11
<b>USDA-ARS Agreement ID:</b>	59-0206-9-075
<b>USDA-ARS Agreement Title:</b>	Mapping and Sequencing of CHR. 2H Bin 10 FHB Resistance QTL for Gene Discovery.
<b>FY11 USDA-ARS Award Amount:</b>	\$ 61,095

**USWBSI Individual Project(s)**

<b>USWBSI Research Category*</b>	<b>Project Title</b>	<b>ARS Award Amount</b>
BAR-CP	Genetic and Physical Mapping of the chr. 2H Bin 10 FHB Resistance QTL.	\$ 61,095
	<b>Total ARS Award Amount</b>	<b>\$ 61,095</b>



Principal Investigator

7/11/12

Date

\* MGMT – FHB Management  
 FSTU – Food Safety, Toxicology, & Utilization of Mycotoxin-contaminated Grain  
 GDER – Gene Discovery & Engineering Resistance  
 PBG – Pathogen Biology & Genetics  
 BAR-CP – Barley Coordinated Project  
 DUR-CP – Durum Coordinated Project  
 HWW-CP – Hard Winter Wheat Coordinated Project  
 VDHR – Variety Development & Uniform Nurseries – Sub categories are below:  
 SPR – Spring Wheat Region  
 NWW – Northern Soft Winter Wheat Region  
 SWW – Southern Soft Red Winter Wheat Region

**Project 1:** *Genetic and Physical Mapping of the chr. 2H Bin 10 FHB Resistance QTL.*

**1. What major problem or issue is being resolved relevant to Fusarium head blight (scab) and how are you resolving it?**

Breeding for resistance to a hemi-biotrophic disease organism, such as Fusarium Head Blight (FHB), is a slow incremental process. To develop new genetic modification approaches, we need to understand the resistance genes and mechanisms involved. There are no single strong resistance genes known for FHB. The Chromosome 2H bin 10 QTL identified in hoCI4196 is one of the strongest FHB resistance QTL known and therefore presents a target for cloning and characterization. Cloning QTL is difficult. In order to facilitate the process we sequenced 36 Bacterial Artificial Chromosome (BAC) clones from the target region in order to identify potential candidate genes and develop improved molecular markers. The candidate genes will be useful as molecular markers in Molecular Marker Assisted Selection (MMAS) and will facilitate identification and validation of resistance QTL from other sources. They should also facilitate understanding of the FHB resistance mechanism.

Line CIho4196 carries good to excellent FHB resistance, but it is a poor breeding parent.

**2. List the most important accomplishment and its impact (i.e. how is it being used) to minimize the threat of Fusarium head blight or to reduce mycotoxins. Complete both sections (repeat sections for each major accomplishment):**

**Accomplishment:**

We sequenced 36 BAC clones from the bin 10 region, 12 from each region. The region 1 sequences were assembled in 110 contigs ranging in size from 84.37 kb to 101 bp. Of the 110 contigs, 32 were 10 kb or more representing a total sequence of 870 kb. The FGENESH gene finder program (<http://linux1.softberry.com>) identified 139 putative genes from region 1. This is the region of bin 10 that we believe is most likely to hold the FHB resistance QTL. Putative gene sequence were blasted against the non-redundant protein sequence database using Blastp resulting in only 22 genes with probable known function. The majority of the remaining sequences were due to probable retrotransposons although there were considerable number of “no hit” sequences. To facilitate further development of the physical map, the gene order developed in this study is being compared to the databases available at IPK-Gatersleben (<http://webblast.ipk-gatersleben.de/barley/viroblast.php>) and Harvest barley genome version 0.05 ([www.harvest-blast.org](http://www.harvest-blast.org)).

Regions 2 and 3 were analyzed in a similar manner

**Impact:**

Sequence data and gene order information will facilitate development of new and better positioned molecular markers to facilitate Molecular Marker based selection of the resistance trait.

**Accomplishment:**

To rapidly facilitate the breeding effort, we identified a mutant that converted the 2-rowed CIho4196 plant to 6-rowed type while retaining the same level of resistance as CIho4196. This line has been widely distributed to breeders. An improved set of primers that facilitate selection of the mutant type *vrs1* gene (6-rowed) were developed. A male sterile mutant was developed and distributed to facilitate crossing. The early and semi-dwarf mutants have not shown good agronomic potential so far.

**Impact:**

The 6-rowed mutant has been widely distributed to barley breeders. The male sterile mutant also has been distributed and should facilitate crossing with the FHB resistant parent. It is expected that these mutants will positively impact the development of 6-rowed FHB resistant cultivars suitable for US barley growers.

**Include below a list of the publications, presentations, peer-reviewed articles, and non-peer reviewed articles written about your work that resulted from all of the projects included in the grant. Please reference each item using an accepted journal format. If you need more space, continue the list on the next page.**

None