High through-put genomics in small grain cereals and applications to molecular breeding

Seungho Cho*, David F. Garvin*† and Gary J. Muehlbauer*

*Department of Agronomy and Plant Genetics, University of Minnesota, St. Paul, MN 55108; and †Plant Science Research Unit, United States Department of Agriculture-Agricultural Research Service, St. Paul, MN 55108

Identification of genes associated with beneficial traits has been one of the major goals in genetics. To facilitate this process in barley genetics, we applied microarray and *in silico* comparative genomics tools. For large-scale physical mapping of barley genes, we used the 22K Barley1 Affymetrix GeneChip to detect barley transcripts in wheat-barley chromosome addition lines harboring barley chromosome2H, 3H, 4H, 5H, 6H, or 7H. In total, 5,543 barley transcripts were detected in the wheat-barley disomic chromosome addition lines, and they were physically mapped to six barley chromosomes based on their barley chromosome-specific detection patterns. Physical locations of these barley transcripts on each chromosome were validated by genomic PCR and *in silico* comparative mapping against the wheat and rice genomes.

We also used the Barley1 GeneChip to identify genes at the QTL for fungal resistance in barley. Deoxynivalenol (DON) is a trichothecene mycotoxin produced by *Fusarium graminearum* during infection and is an important quality-determining factor in harvested barley grain. Our objective was to identify genes associated with reduced DON accumulation. Using a pair of near-isogenic lines (NILs) carrying segregating alleles only at the QTL region on chromosome 3H associated with deoxynivalenol (DON) accumulation upon *F. graminearum* infection, abundance of 22,792 transcripts and their genetic association with DON accumulation were tested. We identified 7 barley transcripts showing differential abundance between the two NILs. By *in silico* comparative mapping, six genes were found on either barley chromosome 3H or

syntenous wheat or rice chromosomes. Our results show that detection of chromosome- or allele-specific transcripts using GeneChips in combination with *in silico* comparative mapping is an efficient tool to explore large-sized plant genomes to identify candidate genes.

We are currently conducting a similar study in wheat to identify genes for Fusarium head blight resistance located at the QTL on wheat chromosome 3BS. Candidate genes showing differential expression between two NILs carrying either resistant or susceptible alleles at the 3BS QTL were identified by the 66K Wheat Affymetrix GeneChip. In addition to transcriptome analysis in host plants, transcriptome change in F. graminearum through infection process is being investigated using the 18K Fusarium Affymetrix GeneChip. Simultaneous transcriptome investigation in plants and a fungal pathogen will facilitate understanding plant-microbe interaction and identification of defense response genes and potentially disease resistance genes.

SEUNGHO CHO

Present Address: University of Minnesota, Agronomy and Plant Genetics, 1991, Buford Circle, Room 411, Borlaug Hall, St. Paul, MN, 55108

E-mail: choxx048@umn.edu

Education

B.Agr. Department of Forest Resources, College of Natural Resources, Korea University, (1988-1992)

M.Agr. in Forest Genetics and Ecology, Department of Forest Resources, College of Natural Resources, Korea University (1993-1995)

Ph.D. in Genetics and Cell Biology, School of Molecular Biosciences, Washington State University (1999-2003)

Professional Experience

Post-doctoral research associate to Dr. Gary J. Muehlbauer, Agronomy and Plant Genetics, University of Minnesota, St. Paul, MN.

Publications

1. Cho S., Kumar J., Shultz J.L., Anupama K., Tefera F., F.J. Muehlbauer (2002) Mapping genes for double podding and other morphological traits in chickpea. EUPHYTICA 128 (2): 285-292.

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