Expression-QTL mapping in wheat to identify genes involved in resistance to *Fusarium graminearum*

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Abstract

Fusarium head blight (FHB) caused by Fusarium graminearum frequently leads to large yield losses in wheat and other cereals but also reduces quality due to contamination with mycotoxins. Breeding for resistance against the disease in wheat provides the best means to protect yields. In the present study, we aim to identify genes which are differentially expressed in response to Fusarium in wheat. Therefore, we employ 200 doubled haploid lines and the two parents (the resistant line CM82036 and the susceptible European spring wheat cultivar Remus) in an expression quantitative trait loci (eQTL) mapping experiment using microarray technology. Expression QTL starts with generating a genetic map using differentially expressed genes between lines as molecular markers to identify genomic regions involved in Fusarium resistance. Microarrays allow to measure differential transcript abundance by detecting florescent emission of labeled-mRNA hybridized to complementary probe sets. We used a custom-build Agilent-microarray

References

BUERSTMAYR H, STEINER B, HARTL L, GRIESSER M, ANGERER N, LENGAUER D, MIEDANER T, SCHNEIDER B, LEMMENS M, 2003: Molecular mapping of QTLs for Fusarium head blight resistance in spring wheat. II. Resistance to fungal penetration and spread. Theor Appl Genet 107: 503-508. to identify genes differentially regulated under varying experimental conditions (e.g. time, genotype, Fusarium vs. water inoculation). Our array-design allows to detect 44000 wheat genes, several hundred wheat candidate genes that have been reported upregulated in response to Fusarium stress in literature and the entire transcriptome of Fusarium graminearum (ca. 14000 genes). In total, we aim to hybridize about 400 microarrays. Once finished, we are able to compare and correlate the genetic map derived from conventional QTL analysis with the new eQTL map. We expect to identify new QTL involved in Fusarium resistance that encode for resistance genes or regulative hotspots controlling multiple genes not encoded on the eQTL. Such hotspots and the related genes cannot be detected by conventional QTL mapping and enables us to build biochemical pathways that are relevant for resistance against Fusarium.

Keywords

Expression QTL, microarray, QTL analysis, transcript abundance

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