Poster Abstract – A.15

FINE MAPPING OF TWO QTL FOR HETEROSIS IN MAIZE

PEA G.*'**, PAULINE SANDRA P.**, CANÈ M.A.***, HTAY-AUNG H.**, LANDI P.***, MORGANTE M.****, PORCEDDU E.****, PÈ M. E.*'**, FRASCAROLI E.***

*) Department of Biomolecular Sciences and Biotechnology, University of Milan, Via Celoria 26, 20133 Milano 8 (Italy)

**) Sant'Anna School for Advanced Studies of Pisa, P.zza Martiri della Libertà 33, 56127 Pisa (Italy)

***) Department of Agroenvironmental Sciences and Technologies "DiSTA", University of Bologna, Via Fanin 44, 40127 Bologna (Italy)

****) Institute of Applied Genomics, Science and Technology Park "Luigi Danieli", Via Linussio 51, 33100 Udine (Italy)

*****) Department of Agrobiology and Agrochemistry, University of Tuscia, Via De Lellis, 01100 Viterbo (Italy)

heterosis, QTL, maize, residual heterozygous line, near isogenic line

Although heterosis is widely exploited for crop improvement and breeding, a clear understanding of its genetic bases is still elusive. In a previous work undertaken to shed light on the genetic basis of heterosis in maize (*Zea mays* L.), we applied a joint classical genetic and QTL analysis to a population of Recombinant Inbred Lines (RIL- $F_{12:13}$) originated from the single cross B73 x H99. Level of heterosis for several agronomic traits and underlying genetic effects were evaluated, together with the relationship between level of heterozygosity and phenotypic performance, and several QTL with heterotic effects on phenotypes were detected. Based on these findings, we followed an introgression program employing marker-assisted breeding on residual heterozygous lines (RHL) to produce pairs of NILs homozygous either for one or the other parental inbred allele (i.e. B73 or H99) at the selected heterotic QTL regions. Large F_2 populations, each segregating only for the respective QTL region, were produced from F_1 hybrids obtained by crossing contrasting NILs for each QTL.

In this work we describe the results of our research work aimed at the fine mapping of two of the introgressed QTL mapped on chromosome 4 (bin 10) and 10 (bin 03). In particular, 3840 and 1152 F_2 plants were genotyped at markers flanking QTL 4.10 and QTL 10.03 respectively, and F_3 families were produced by selfing F_2 plants recombinant at the respective QTL region. The so obtained F_3 families are highly-informative and will be evaluated for agronomic traits in order to fine map their respective QTL.