

**Supplementary file 3.** Methodological specifications of the two approaches used to link genotypes and phenotypes

	<b>Linkage analysis</b>	<b>Linkage disequilibrium mapping</b>
<b>Population type</b>	Biparental cross with known pedigree: $Y=\mu+M+e$	Population with unknown pedigree/structure: $Y=\mu+M+Q+K+e$
<b>Nb of segregating alleles</b>	LOW	HIGH
<b>Resolution</b> (i.e. nb of meiotic recombination events that have occurred between a DNA marker and a QTL)	LOW Few recombination between markers within the studied pedigree	HIGH Historical recombination within the unobserved pedigree from which the studied population was derived
<b>Nb of markers</b> needed to detect a QTL	LOW (low-density linkage maps)	- Extremely HIGH for genome-wide investigations - Targeted candidate genes
<b>QTL confidence interval</b>	LARGE (cM) Loose marker-trait association	NARROW (depends on LD window) Tight marker-trait association down to the causal variant