**Supplementary Data S1. R package LPmerge Script.**

library(LPmerge)

library(Rglpk)

setwd("C:/Users/Du/Desktop/mungbean/")

map.names = c("H", "I", "W1", "L", "W2", "W3", "Y")

#############1

map.names = c("H", "I", "W1", "L", "W2", "W3", "Y")##1

Maps <- list()

i <- 1

for (i in 1:7) {

filename <- paste(map.names[i],".csv",sep="")

 input <- read.csv(filename,header=T,as.is=T,check.names=F)

 Maps[[i]] <- input[which(input$chr=="8"), c(1,3)]

}

names(Maps) <- map.names

str(Maps)

print(link.map.lengths <- unlist(lapply(Maps,function(x){max(x$cM)})))

mean(link.map.lengths)

unweighted <- LPmerge(Maps,max.interval=1:4)

head(unweighted[[1]])

write.csv(unweighted[[1]], file = "CA\_8.txt")

sink("out\_CA\_8.txt")

print(input$marker)

print(input$chr)

print(input$cM)

print("link.map.lengths results")

print("map.names results")

print("unweighted <- LPmerge(Maps,max.interval=1:4)")

print(unweighted)

sink()

**Supplementary Data S2. The optimal number of MQTL per chromosome was determined based on the model selection criteria (\_model.txt) as well as the 95% confidence intervals (CIs) and peak locations of the MQTL (\_table.txt).**

LG1

\_model.txt



\_table.txt



LG2

\_model.txt



\_table.txt



LG3

\_model.txt



\_table.txt



LG4

\_model.txt



\_table.txt



LG5

\_model.txt



\_table.txt



LG6

\_model.txt



\_table.txt



LG7

\_model.txt



\_table.txt



LG8

\_model.txt



\_table.txt



LG9

\_model.txt



\_table.txt



LG10

\_model.txt



\_table.txt



LG11

\_model.txt



\_table.txt

