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### rPlasmo tools, Tools for viewing and analyzing malaria data.
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#####
##          Code for computing mQTL
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## Read input files
markerList <- readMark() ## QTL_genotype.csv file
pheno <- readPeak() ## QTL_phenotype.csv file
covar <- c(1, 2, 5, 6, 5, 1, 6, 1, 6, 2, 2, 4, 2, 5, 3, 4, 1, 5, 1,
       6, 2, 2, 4, 3, 1, 5, 3, 3, 1, 4, 3, 1, 3, 4, 3)

## Compute QTLs
final <- mzQTL(pheno = pheno, markerList = markerList, covar = covar)
mzAlpha <- c( 0.05, 0.01, 0.001, 0.05 / (nrow(pheno) - 2) )
alphaT <- matrix( NA, nrow = nrow(pheno), ncol = length(mzAlpha) )
for( i in 1:length(mzAlpha) ){
  alphaT[,i] <- sapply(final[[2]], function(x) summary(x, alpha =
mzAlpha[i]) )
}
alphaT <- data.frame(alphaT)
names(alphaT) <- mzAlpha
row.names(alphaT) <- pheno$compound
chr <- as.numeric(as.vector(unlist(sapply(final[[1]],
  function(x) max(x)[1]))))
pos <- as.numeric(as.vector(unlist(sapply(final[[1]],
  function(x) max(x)[2]))))
lod <- as.numeric(as.vector(unlist(sapply(final[[1]],
  function(x) max(x)[3]))))
pVal <- rep(NA, length(lod))
for( i in 1:length(final[[1]]))
  pVal[i] <- min(summary(final[[1]][[i]], perms = final[[2]][[i]], pvalues =
T)$pval)

## Generate summary
cpdSummary <- data.frame(
  compound = names(final[[1]]),
  chr, pos, lod, pVal, stringsAsFactors = F)
cpdSummary <- cbind(cpdSummary, alphaT)

#####
##          Code for modeling parasite growth
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#######
## Read in observed competition data and fitting parameters
dat <- readPeak() ## population_summary.csv file

## Generate 50,000 random sampling points for grid search
fitR <- fitComp( obs = dat$S_1, pop1 = .5, syncC = F, N = 50000)

## Final fit used for Figure 6A (50:50 population)
fitF1 <- fitComp(obs = dat$S_1,
                  pop1 = .5,
                  lc1 = 47,
                  lc2 = 49,
                  t0 = 43,
                  r1 = .065,
                  sc = 8,
                  pSync = 1,
                  rSample = F,
                  syncC = T
)
## Final fit used for Figure 6B (25:75 population)
fitF2 <- fitComp(obs = dat$S_2,
                  pop1 = .25,
                  lc1 = 47,
                  lc2 = 49,
                  t0 = 43,
                  r1 = .065,
                  sc = 8,
                  pSync = 1,
                  rSample = F,
                  syncC = T
)

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