

1. why study multiple traits together?

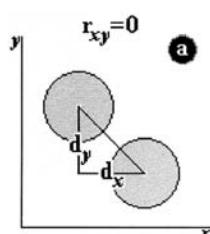
- avoid reductionist approach to biology
 - address physiological/biochemical mechanisms
 - Schmalhausen (1942); Falconer (1952)
- separate close linkage from pleiotropy
 - 1 locus or 2 linked loci?
- identify epistatic interaction or canalization
 - influence of genetic background
- establish QTL x environment interactions
- decompose genetic correlation among traits
- increase power to detect QTL

QTL II: GxE

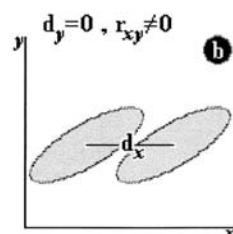
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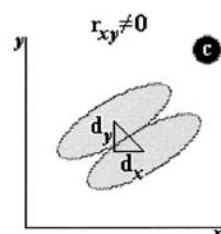
interplay of pleiotropy & correlation



pleiotropy only



correlation only



both

Korol et al. (2001)

QTL II: GxE

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3 correlated traits (Jiang Zeng 1995)

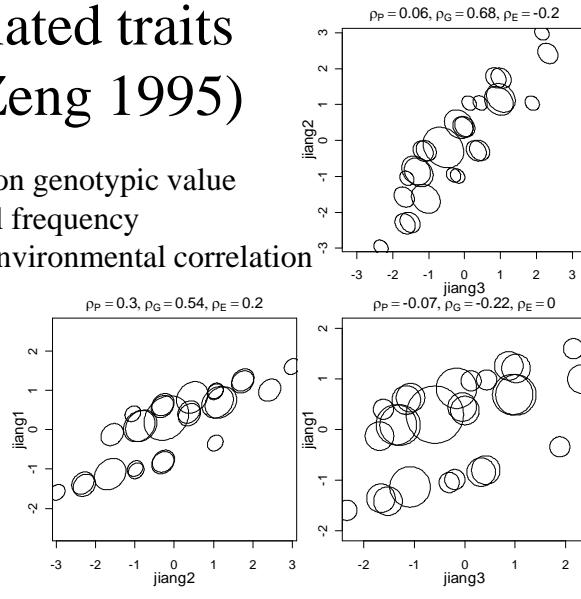
ellipses centered on genotypic value
width for nominal frequency
main axis angle environmental correlation

3 QTL, F2

27 genotypes

note signs of
genetic and
environmental
correlation

QTL II: GxE

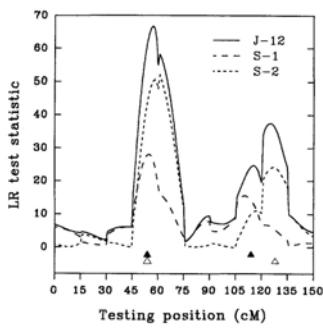


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pleiotropy or close linkage?

2 traits, 2 qtl/trait
pleiotropy @ 54cM
linkage @ 114,128cM
Jiang Zeng (1995)



QTL II: GxE

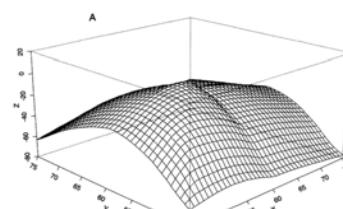
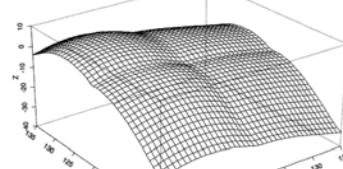


FIGURE 2.—Three-dimensional likelihood surfaces (expressed as deviations from the maximum likelihood, with the peaks on the diagonal) for the test of pleiotropy vs. close linkage. Plot A shows the region between 45 and 75 cM of Figure 1(A) and the region between 105 and 135 cM (B). X is the testing position for the testing trait 1 and Y is the testing position for a QTL affecting trait 2. In both plots, two QTLs are located in the same position and simultaneously are tested. In plot A, pleiotropic QTLs are located in the same place.



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QTL with GxE or Covariates

- adjust phenotype by covariate
 - covariate(s) = environment(s) or other trait(s)
- additive covariate
 - covariate adjustment same across genotypes
 - “usual” analysis of covariance (ANCOVA)
- interacting covariate
 - address GxE
 - capture genotype-specific relationship among traits
- another way to think of multiple trait analysis
 - examine single phenotype adjusted for others

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R/qtl & covariates

- additive and/or interacting covariates
 - test for QTL after adjusting for covariates
- ```
Get Brassica data.
library(qtlbim)
data(Bnapus)
Bnapus <- calc.genoprob(Bnapus, step = 2, error = 0.01)

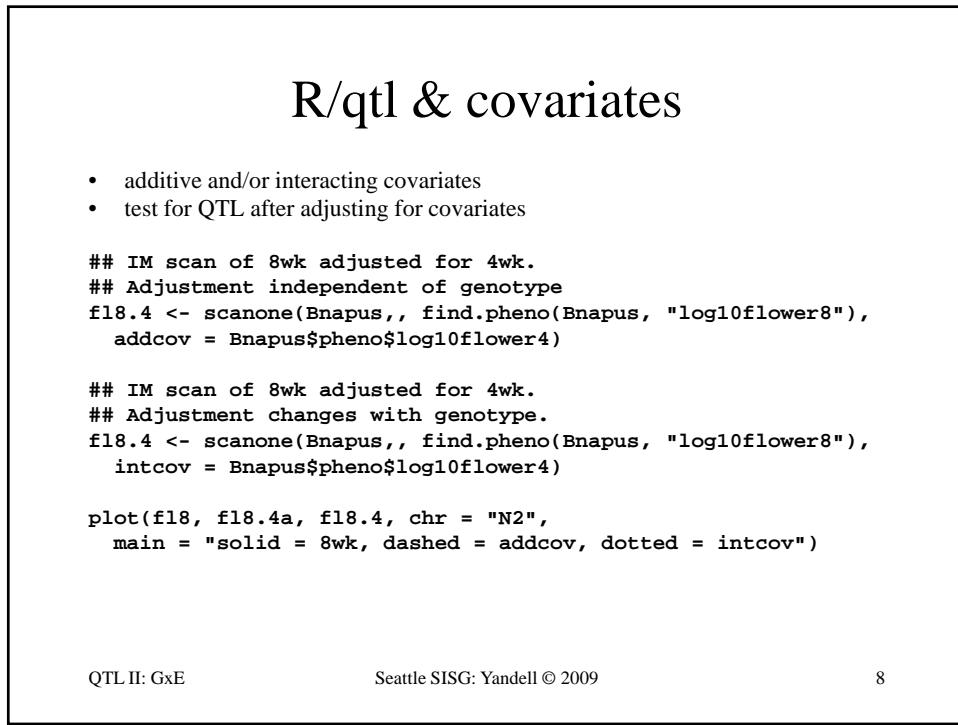
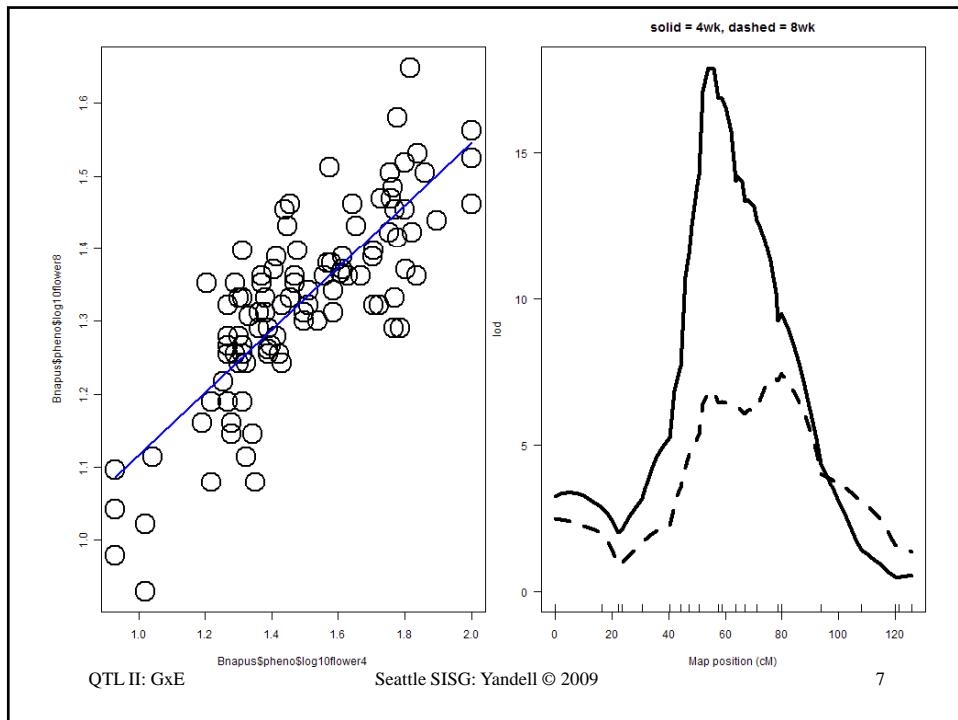
Scatterplot of two phenotypes: 4wk & 8wk flower time.
plot(Bnapus$pheno$log10flower4,Bnapus$pheno$log10flower8)

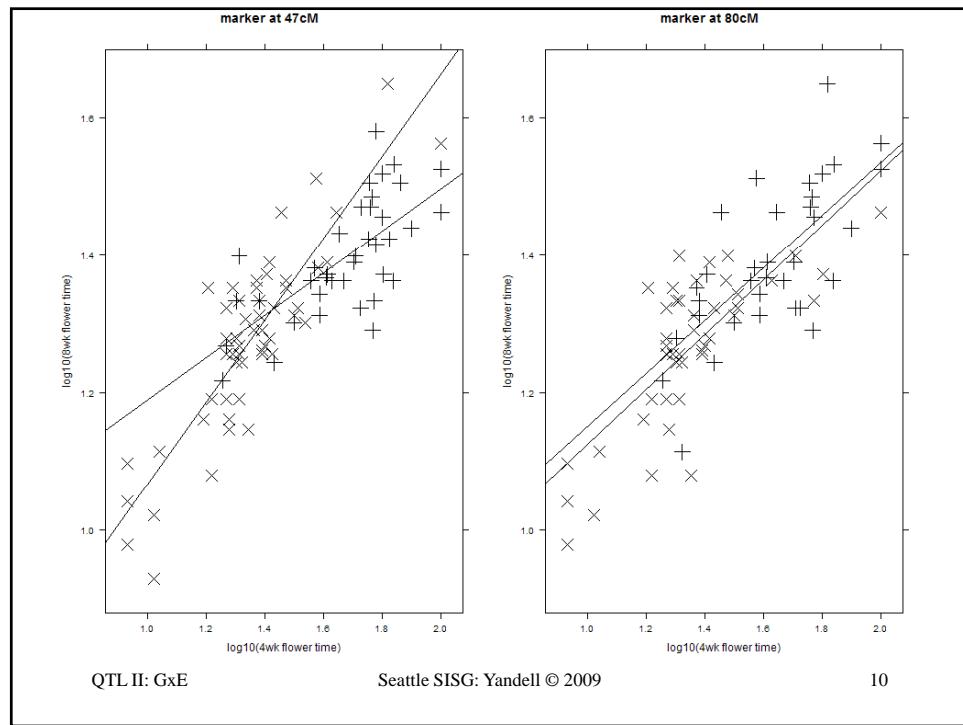
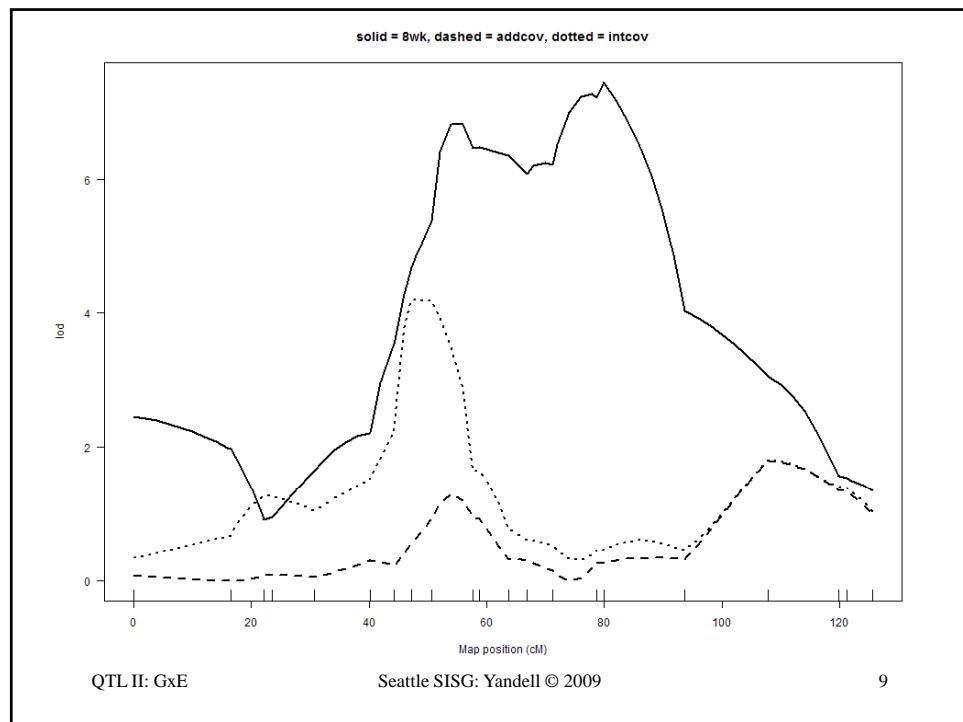
Unadjusted IM scans of each phenotype.
f18 <- scanone(Bnapus,, find.pheno(Bnapus, "log10flower8"))
f14 <- scanone(Bnapus,, find.pheno(Bnapus, "log10flower4"))
plot(f14, f18, chr = "N2", col = rep(1,2), lty = 1:2,
 main = "solid = 4wk, dashed = 8wk", lwd = 4)
```

QTL II: GxE

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## scatterplot adjusted for covariate

```
Set up data frame with peak markers, traits.
markers <- c("E38M50.133","ec2e5a","wg7f3a")
tmpdata <- data.frame(pull.genotype(Bnapus)[,markers])
tmpdata$f14 <- Bnapus$pheno$log10flower4
tmpdata$f18 <- Bnapus$pheno$log10flower8

Scatterplots grouped by marker.
library(lattice)
xyplot(f18 ~ f14, tmpdata, group = wg7f3a,
 col = "black", pch = 3:4, cex = 2, type = c("p","r"),
 xlab = "log10(4wk flower time)",
 ylab = "log10(8wk flower time)",
 main = "marker at 47cM")
xyplot(f18 ~ f14, tmpdata, group = E38M50.133,
 col = "black", pch = 3:4, cex = 2, type = c("p","r"),
 xlab = "log10(4wk flower time)",
 ylab = "log10(8wk flower time)",
 main = "marker at 80cM")
```

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## R/qtlbim and GxE

- similar idea to R/qtl
  - fixed and random additive covariates
  - GxE with fixed covariate
- multiple trait analysis tools coming soon
  - theory & code mostly in place
  - properties under study
  - expect in R/qtlbim later this year
  - Samprit Banerjee (N Yi, advisor)

QTL II: GxE

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