

Prototype QTL Strategy: Phenotype bp in Cross hyper

Brian S. Yandell, W. Whipple Neely, Nengjun Yi

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Overview

Initialization

1-D & 2-D Scans

Anova Fit

User Customized Section

Conclusion

Automated Strategy

- ▶ Estimate positions and effects of main QTL.
- ▶ Find chromosomes with epistasis.
- ▶ Estimate epistatic pair positions and effects.
- ▶ Confirm genetic architecture with ANOVA.

Running Sweave

```
> library(qtlbim)

> qb.sweave(hyper, pheno.col = 1,
+ n.iter = 3000, n.draws = 8,
+ scan.type = "2logBF", hpd.level = 0.5,
+ threshold = c(upper = 2),
+ SweaveFile = "",
+ SweaveExtra = "/tmp/Rinst1061764704/qtlbim/external/hyper.slide.extra.Rnw",
+ PDFDir = "bpPDF",
+ remove.qb = TRUE)
```

Cross Object

```
> summary(cross)
```

Backcross

No. individuals: 250

No. phenotypes: 2

Percent phenotyped: 100 100

No. chromosomes: 19

Autosomes: 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19

Total markers: 170

No. markers: 22 8 6 20 14 11 7 6 5 5 14 5 5 5 11 6 12 4 4

Percent genotyped: 47.9

Genotypes (%): BB:50.1 BA:49.9

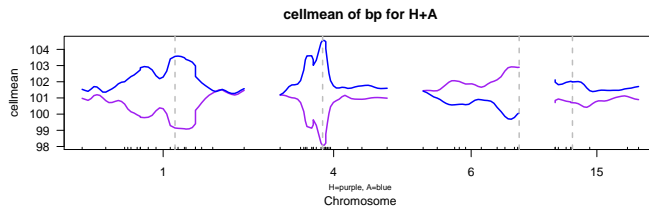
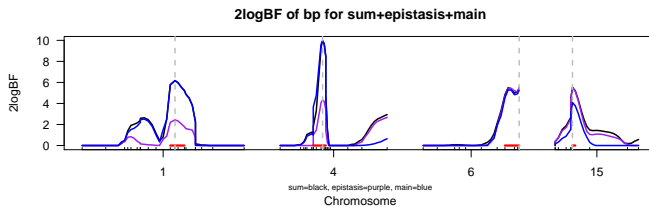
Create MCMC runs

```
> cross <- qb.genoprob(cross,step=2)
> cross.qb <- qb.mcmc(cross, pheno.col = pheno.col,
+   genoupdate=TRUE, n.iter = 3000, verbose=FALSE)
```

1-D 2logBF Scan

```
> hpd.level  
[1] 0.5  
  
> scan.type  
[1] "2logBF"  
  
> cross.hpd <- qb.hpdone(cross.qb, hpd.level, scan.type)  
> sum.one <- summary(cross.hpd)  
> sum.one  
  
chr n.qtl pos lo.50. hi.50. 2logBF A H  
1 1 0.826 67.8 64.5 74.3 6.181 103.568 99.143  
4 4 3.035 29.5 23.0 31.7 9.970 104.552 98.073  
6 6 0.873 66.7 56.8 66.7 5.488 99.710 102.866  
15 15 0.395 17.5 17.5 19.5 5.448 101.993 100.703  
  
> chrs <- as.vector(sum.one[, "chr"])  
> pos <- sum.one[, "pos"]  
  
> plot(cross.hpd)
```

1-D Scan: 2logBF Profile



2-D: find epistatic pairs

```
> two <- qb.scantwo(cross.qb, chr = chrs, type = scan.type)
> sum.two <- summary(two, sort = "upper", threshold = threshold,
+   refine = TRUE)
> sum.two
```

upper: 2logBF of bp for epistasis
 lower: 2logBF of bp for full
 Thresholds: upper=2

	n.qtl	l.pos1	l.pos2	lower	u.pos1	u.pos2	upper
c6 :c15	1.004	59.0	17.5	11.89	59.0	17.5	11.90
c4 :c6	1.185	29.5	59.0	13.77	74.3	61.2	7.49
c4 :c15	1.452	29.5	17.5	13.61	74.3	47.6	6.84
c15:c15	0.261	17.5	33.5	7.54	17.5	31.5	6.52
c1 :c4	1.817	67.8	29.5	14.41	72.1	29.5	6.10
c1 :c15	1.255	67.8	17.5	11.20	72.1	17.5	5.23
c1 :c6	1.103	67.8	59.0	11.37	67.8	59.0	5.21
c1 :c1	0.366	43.7	77.6	7.46	39.4	77.6	5.20
c4 :c4	0.417	29.5	74.3	11.00	28.4	49.5	4.76
c6 :c6	0.111	61.2	65.6	7.52	40.4	56.8	3.94

Initial Genetic Architecture

```
> cross.arch <- qb.arch(sum.two, chrs, pos)
> cross.arch
```

main QTL loci:

```
      [,1]      [,2]      [,3]      [,4]      [,5]      [,6]      [,7]      [,8]      [,9]
chr  "1"      "1"      "15"     "15"     "15"     "4"      "4"      "4"      "6"
pos  "39.35"  "71.48"  "17.50"  "31.52"  "47.64"  "29.13"  "49.45"  "74.30"  "40.40"
      [,10]
chr  "6"
pos  "60.54"
```

Epistatic pairs by qtl, chr, pos:

	qtl	qtlb	chra	chrb	posa	posb
pair 1	3	10	15	6	17.50	60.54
pair 2	8	10	4	6	74.30	60.54
pair 3	5	8	15	4	47.64	74.30
pair 4	3	4	15	15	17.50	31.52
pair 5	2	6	1	4	71.48	29.13
pair 6	2	3	1	15	71.48	17.50
pair 7	2	10	1	6	71.48	60.54
pair 8	1	2	1	1	39.35	71.48
pair 9	6	7	4	4	29.13	49.45
pair 10	9	10	6	6	40.40	60.54

Epistatic chromosomes by connected sets:

1,15,4,6

Construct QTL Object

use R/qtl tools to check model fit
first simulate missing markers
then construct QTL object

```
> cross.sub <- subset(cross, chr = unique(cross.arch$qtl$chr))  
> n.draws  
  
[1] 8  
  
> cross.sub <- sim.geno(cross.sub, n.draws = n.draws, step = 2,  
+   error = 0.01)  
> qtl <- makeqtl(cross.sub, as.character(cross.arch$qtl$chr), cross.arch$qtl$pos)  
> cross.sub <- clean(cross.sub)
```

Stepwise Reduction

```
> cross.step <- step.fitqtl(cross.sub, qtl, pheno.col, cross.arch)
```

	drop	LOD	p
1	Chr1@39.3:Chr1@71.3	-0.04690	1.0000
2	Chr4@29.5:Chr4@50	0.03090	0.7180
3	Chr4@29.5:Chr4@50	0.03410	0.7030
4	Chr4@29.5:Chr4@50	0.04260	0.6700
5	Chr4@29.5:Chr4@50	0.04320	0.6670
6	Chr4@29.5:Chr4@50	0.05010	0.6430
7	Chr15@31.5	-0.12600	1.0000
8	Chr4@29.5:Chr4@50	0.07130	0.5780
9	Chr6@40.4:Chr6@60	0.00674	0.8640
10	Chr4@29.5:Chr4@50	0.05790	0.6150
11	Chr4@29.5:Chr4@50	0.07510	0.5660
12	Chr15@47.5	0.12300	0.4610
13	Chr4@50	0.14900	0.4170
14	Chr6@40.4	0.09380	0.5180
15	Chr4@74.3	0.84400	0.0521

```
> summary(cross.step$fit)
```

	df	SS	MS	LOD	%var	Pvalue(Chi2)	Pvalue(F)
Model	5	4844.318	968.86365	17.39575	27.41715	7.771561e-16	1.554312e-15
Error	244	12824.618	52.55991				
Total	249	17668.936					

Stepwise Reduction

	df	Type III SS	LOD	%var	F value	Pvalue(F)	
Chr1@39.3	1	290.212	1.215	1.642	5.522	0.019581	*
Chr1@71.3	1	656.678	2.711	3.717	12.494	0.000488	***
Chr15@17.5	1	323.724	1.353	1.832	6.159	0.013748	*
Chr4@29.5	1	2747.049	10.536	15.547	52.265	6.21e-12	***
Chr6@60	1	473.140	1.967	2.678	9.002	0.002977	**

Reduced Genetic architecture

```
> cross.arch <- cross.step$arch  
> cross.arch
```

main QTL loci:

	1	2	3	6	10
chr	"1"	"1"	"15"	"4"	"6"
pos	"39.35"	"71.48"	"17.50"	"29.13"	"60.54"

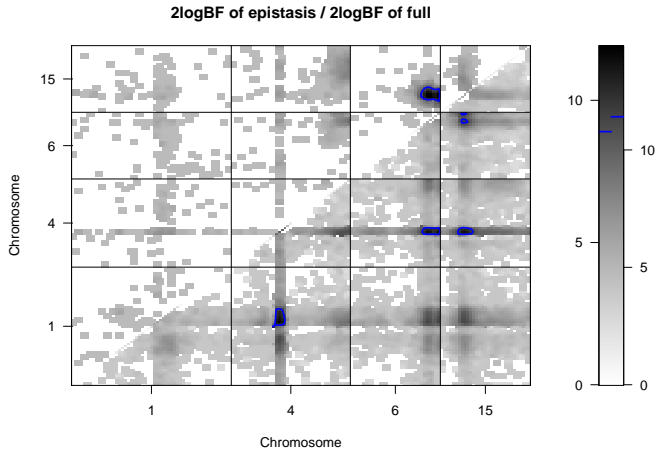
epistatic pairs: none

2-D Plots

2-D plots by cliques (if any epistasis)

```
> for(i in names(cross.arch$chr.by.set))  
+   plot(two, chr = cross.arch$chr.by.set[[i]], smooth = 3,  
+       col = "gray", contour = 3)
```

2-D Plots: clique 1

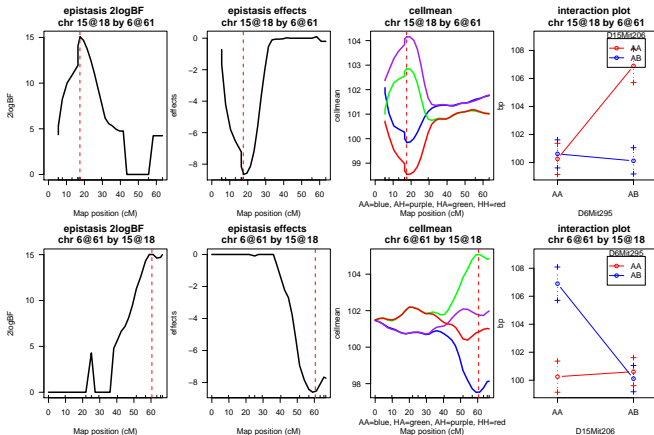


Slice Each Epistatic Pair

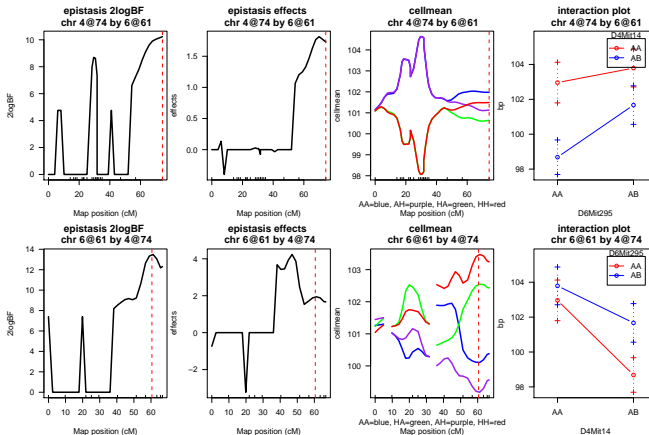
show detail plots for epistatic pairs (if any)

```
> if(!is.null(cross.arch$pair.by.chr)) {  
+   for(i in seq(nrow(cross.arch$pair.by.chr$chr))) {  
+     chri <- cross.arch$pair.by.chr$chr[i,]  
+     posi <- cross.arch$pair.by.chr$pos[i,]  
+     if(chri[1] != chri[2])  
+       plot(qb.slicetwo(cross.qb, chri, posi, scan.type))  
+   }  
+}
```

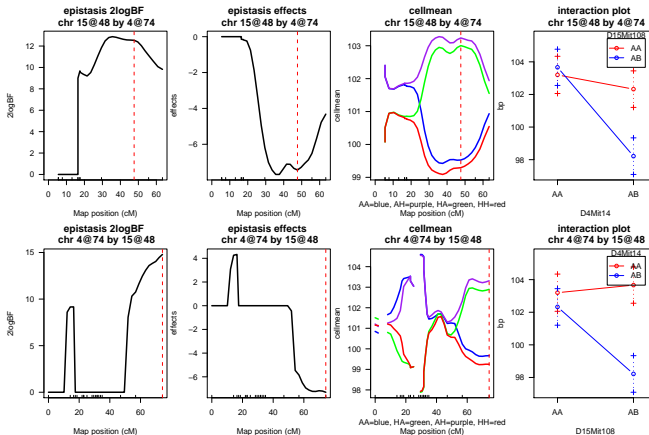
Epistatic Pair 15 and 6



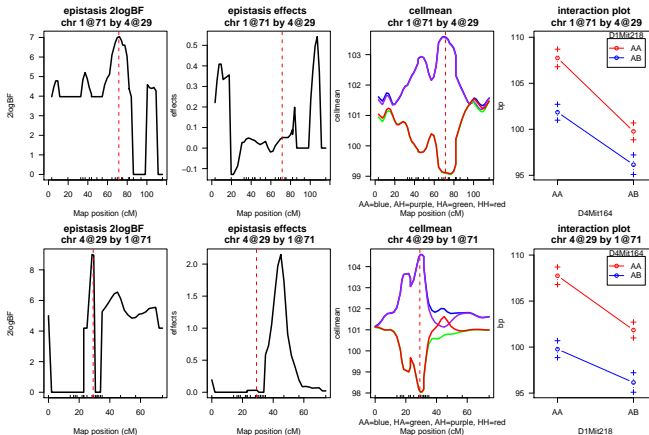
Epistatic Pair 4 and 6



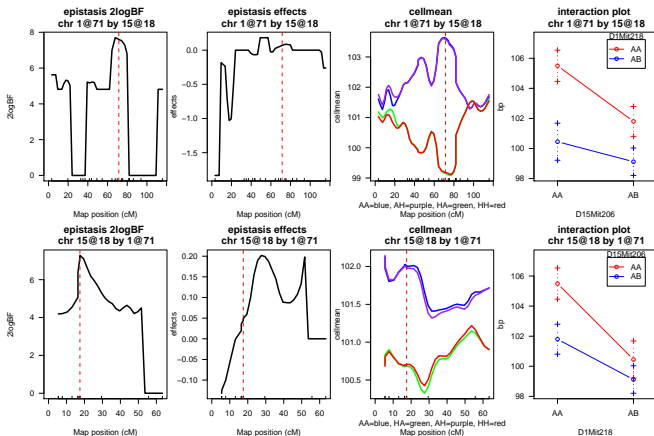
Epistatic Pair 15 and 4



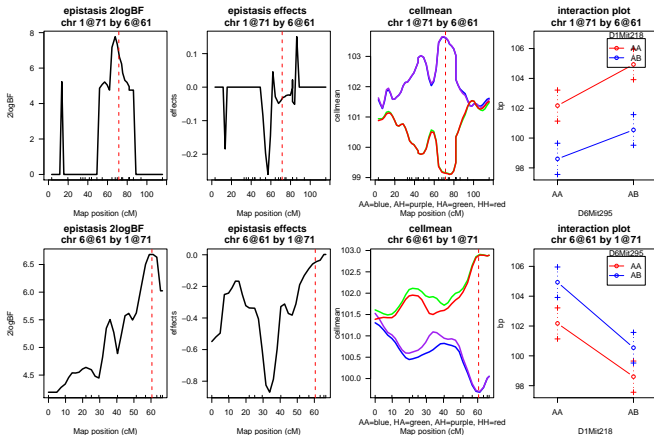
Epistatic Pair 1 and 4



Epistatic Pair 1 and 15



Epistatic Pair 1 and 6



Compare with Literature

Sugiyama et al. (2002) found:
two main QTLs on 1 4
two epistatic pairs with 6.15, 7.15
compare to present model:

```
> arch3 <- qb.arch(cross.step, main = c(1, 4), epistasis = data.frame(q1 = c(6,  
+ 7), q2 = rep(15, 2)))  
> arch3
```


Sugiyama Model

```
> cross.step2 <- step.fitqtl(cross.sub, qtl, pheno.col, arch3)
> summary(cross.step2$fit)
```

Sugiyama vs. Automata

formal comparison with automated model

```
> anova(cross.step, cross.step2)
```

final tasks:

externally rename file .tex to bp.tex

and run pdflatex twice on it

remove objects created by R/qt1bim if desired

```
> file.rename(".tex", "bp.tex")
> invisible(system("pdflatex bp.tex",intern=TRUE))
> invisible(system("pdflatex bp.tex",intern=TRUE))

> remove.qb

[1] FALSE

> if (remove.qb) {
+   qb.remove(cross.qb)
+   rm(cross, cross.sub, pheno.col, threshold, n.iter, n.draws,
+       remove.qb)
+ }
```