# Linkage analysis of the XVIth QTLMAS simulated dataset using QTLMAP 

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## Linkage analysis of the XVI ${ }^{\text {th }}$ QTLMAS simulated dataset using QTLMap

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## Aim

More and more association analyses, often without any use of linkage
$\longrightarrow$ Does LA analyses still add some information?
$\Longrightarrow$ Is LA better than association in false positive discovery?

It could be an interesting comparison point for the organizers

## Dataset



5 chromosomes / 10000 SNP (1/0.05Mb) / 3 traits

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## QTLMap

A software for trait dissection in mixture of full and half-sib families using genomic markers

Single QTL LA Multi QTL<br>LD<br>LDLA

Multi Trait X LA

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## QTLMap

## A software for trait dissection in mixture of full and half-sib families using genomic markers

Single QTL<br>Multi QTL

## LA <br> X LD <br> LDLA

Multi Trait X LA

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## Linkage Analysis

- Full Mixture model, a likelihood approach

Parental phase probability
Likelihood $=\Pi_{i} \Sigma_{h s i} p\left(h s_{i} / M\right) \Pi_{j} \overleftarrow{\Sigma_{h d j} p\left(h d_{j} / h s_{i}, M\right)}$

$$
\Pi_{k} \Sigma_{i j k} p\left(t_{i j k} / h d_{j j} h s_{i j}, M\right) \cdot f\left(y_{i j k} / \tau_{i j k}, \sigma^{2}\right)
$$

Transmission probability at position $x$

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## Linkage Analysis

- linearized model, the regression approach

-The penetrance is a normal with as a mean the weighted QTL effect

$$
y \approx \mathscr{N}\left(\mu, V=I \sigma^{2}\right)
$$

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## Linkage Analysis

- mixed model (including an animal effect)

$$
y \approx \mathscr{N}\left(\mu, V=A \sigma_{a}^{2}+I \sigma_{e}^{2}\right)
$$

> The user gives the pedigree and heritability
> QTLMap computes the relationship matrix A
$>$ The effects are estimated by a BLUP

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## LA vs. LA Animal



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## LA vs. LA Animal

- No difference in QTL detection was observed in this dataset
- It would be interesting to test this method in a small family structured design
$\longrightarrow$ Focus on LA analysis
Use of all the design (G0-G1-G2-G3)
$\longrightarrow 60$ independant families (3000 phenotypes)

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## Single QTL

Detection of 9 QTL using 3000 animals:

- Two were not observed using only the G3 animals
- Surprisingly QTL affecing trait 2 on chromosome 5 is less significant
- A few QTL have a location shift

There are more than one QTL location on Chr 1 and 5
Chr1: 19 (trait 2), 52 (trait 1) and 80cM (trait 3)
Chr5: 71 (trait 2) and 79cM (trait 1)

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## Single QTL

Detection of 9 QTL using 3000 animals:

- Two were not observed using only the G3 animals
- Surprisingly QTL affecing trait 2 on chromosome 5 is less significant
- A few QTL have a location shift
$\longrightarrow$ There are more than one QTL location on Chr 1 and 5

Some of the QTL affecting different traits share the same location (pleiotropy ?)

Chr4: 23 and 26cM for traits 1 and 2, respectively

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## Multi-QTL

0 vs 2 QTL
-No significant QTL

## 1 vs 2 QTL

- Chr1 / Trait 3: 60 and 84cM
- Chr5 / Trait 2: 25 and 68cM

Chr1 seems to carry 3 QTL, around 20, 60 and 84 cM Chr5 seems to carry a QTL at 25 cM in addition of the QTL detected around 70 and 80 cM

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## Multi-Trait

Chromosome 1:
Single- and Multi-QTL analyses:
Chr1 seems to carry 3 QTL, around 20, 60 and 84 cM

Multi-trait analyses:

Single-QTL (three traits):
Single-QTL (traits 1 / 2): Single-QTL (traits 1 / 3): 3QTL (traits $1 / 2 / 3$ ) :

1 QTL at 60cM
1 QTL at 19 cM
1 QTL at 78 cM
QTL at 19, 70 and 80 cM

QTL at 60cM is probably a false positive There are 2 pleiotropic QTL at 19 and 78cM

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## Multi-Trait

Chromosome 4: Multi-trait using traits 1 and 2:

1 QTL at 23.5 cM

The QTL at 23.5 cM is pleiotropic

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## The off topic slide

With the Linkage Disequilibrium (LD) analysis, we do not find the QTL on Chromosome 2 while the QTL on Chr 3 is shifted

## Chr2

LA
LD
0.79 ***
0.80 NS

Chr3
0.36***
0.49***

Using the LA seems to be interesting for Chr2. Hard to conclude for Chr3

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# Conclusion 

Chromosome Trait Location (M) Effect (2a) Effect (trait std)


- 8 QTL were detected, three of them being pleiotropic
- Using different models allows the identification of additional QTL but also to precise locations or avoid false positive
- LA sometimes is more interesting than LD

We will see tomorrow how wrong we are !

## Thank you for your attention



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