

# Markov Chain Monte Carlo Model Composition Search Strategy for Quantitative Trait Loci in a Bayesian Hierarchical Model

Susan J. Simmons, Fang Fang, Qijun Fang, and Karl Ricanek

**Abstract**—Quantitative trait loci (QTL) experiments have yielded important biological and biochemical information necessary for understanding the relationship between genetic markers and quantitative traits. For many years, most QTL algorithms only allowed one observation per genotype. Recently, there has been an increasing demand for QTL algorithms that can accommodate more than one observation per genotypic distribution. The Bayesian hierarchical model is very flexible and can easily incorporate this information into the model. Herein a methodology is presented that uses a Bayesian hierarchical model to capture the complexity of the data. Furthermore, the Markov chain Monte Carlo model composition (MC<sup>3</sup>) algorithm is used to search and identify important markers. An extensive simulation study illustrates that the method captures the true QTL, even under nonnormal noise and up to 6 QTL.

**Keywords**—Bayesian hierarchical model, Markov chain Monte Carlo model composition, quantitative trait loci.

## I. INTRODUCTION

QUANTITATIVE trait loci (QTL) experiments have yielded important biological and biochemical information necessary for understanding the relationship between genetic markers and quantitative traits [1]. There are many examples where the identification of QTL have made substantial impact in industry such as the genetic markers responsible for weight gain in pigs [2], pecking-related traits in chickens [3], and starch content and composition in maize [4].

Over the last 20 years, there has been an abundance of algorithms proposed for QTL mapping such as interval mapping strategies [5]-[8], composite interval mapping strategies [9]-[12], multiple interval mapping [13]-[20], Bayesian interval mapping [21]-[24], and model selection strategies [25]-[27]. However, most algorithms allow only one observation per genotypic distribution. In situations

such as plant QTL experiments where there can be cloned plants, observations within lines are summarized into a single observation. By summarizing the cloned plants into one observation, important information about variation within each line is lost.

In this paper, an MC<sup>3</sup> model selection strategy is proposed in a flexible Bayesian hierarchical model that incorporates replicate information and thereby taking full advantage of variability within a (plant) line. The proposed method is validated through an extensive simulation study involving a variety of QTL scenarios.

## II. METHODOLOGY

### A. Bayesian Hierarchical Model

The measured trait in the QTL experiment is represented by  $y_{ij}$  where  $i=1, \dots, L$  ( $L$  = number of lines) and  $j=1, \dots, n_i$  ( $n_i$  = the number of replicates within line  $i$ ). We assume that the observed trait follows a normal distribution with mean  $\theta_i$  and variance  $\sigma_i^2$  or in other words

$$y_{ij} | \theta_i, \sigma_i^2 \sim N(\theta_i, \sigma_i^2) \quad i = 1, \dots, L. \quad (1)$$

Furthermore, the means are assumed to be influenced by the marker information matrix, which is denoted as  $X$  where  $X$  is  $M \times L$  and  $M$  is the number of markers. The mean of each line is assumed to follow a normal distribution with mean  $X_i' \beta$  and variance  $\tau^2$  where  $X_i'$  is the transpose of  $i^{\text{th}}$  column of  $X$  or

$$\theta_i | X, \beta, \tau^2 \sim N(X_i' \beta, \tau^2). \quad (2)$$

Since no prior information is assumed to be known about which markers might be the QTL, we assign a normal distribution with mean 0 and a large variance of 100 on the coefficient for each marker ( $\beta \sim N(0, 100)$ ).

The prior distribution for the variance parameters  $\sigma_i^2$  and  $\tau^2$  are assigned Inverse- $\chi^2(1)$  which has an infinite mean and variance.

These assumptions yield an implicit full posterior distribution; however, the full conditional distributions have a nice parametric form. The full conditional posterior distributions are represented below as  $\varphi_i$  for random variable  $\varphi$  given all other quantities.

S.J. Simmons is with the Department of Mathematics and Statistics at the University of North Carolina Wilmington, Wilmington, NC 28403 USA (phone: 910-962-3296; fax: 910-962-7107; e-mail: simmonssj@uncw.edu).

F. Fang and Q. Fang were with the Department of Mathematics and Statistics at University of North Carolina Wilmington, Wilmington, NC 28403 USA. They are now with the Department of Mathematics at University of Arizona, Tucson, AZ 85721 USA (e-mail: fangfang1101@gmail.com and fangqijun33@hotmail.com).

K. Ricanek is with the Department of Computer Science at University of North Carolina Wilmington, Wilmington, NC 28403 USA (e-mail: ricanekk@uncw.edu).

The transitional probability  $\alpha_{ij}$  defines the likelihood of the chain moving to the new model,  $j$ . A Bernoulli random variable is generated with probability of success  $\alpha_{ij}$ . If the generated Bernoulli random variable is 1, then the new model,  $j$ , becomes the current model. If, however, the generated Bernoulli random variable is 0, the current model,  $i$ , is maintained as the current model. The algorithm continues by randomly identifying a new position along the current model vector and switching its value. The likelihood of this new model is calculated and a new transitional probability is

where  $a_j$  is the effect of the  $j^{\text{th}}$  marker,  $x_{ij}$  is the value of the  $j^{\text{th}}$  marker of the  $i^{\text{th}}$  line, and  $\varepsilon_{ij}$  is the random noise. The

algorithm defined in Section II was developed under the assumption of normal noise, but error distributions are generally unknown. In this simulation study, it was of interest to determine if this algorithm was robust to the assumption of normality, and therefore, the error distribution simulated in this work was generated from a gamma distribution.

The gamma distribution is a two-parameter, continuous probability distribution controlled by its shape,  $\alpha$ , and scale,  $\phi$ , parameters. We used two different gamma distributions in this study. The first gamma distribution had a shape parameter of 0.5 and a scale parameter of 1, giving an expectation for the variance within each line of 0.5, i.e. a tight variance. The second gamma distribution had a shape parameter of 1 and a scale parameter of 3 giving an expected variance of 3 within each line, i.e. a larger variance for the random noise. To further stress the system, effect sizes,  $a_j$ , were selected that ranged from 1 to 9 where an effect size of 1 is very small, effect size of 5 is moderate, and effect size of 9 is large. In order to adequately evaluate the system for complex QTL marker identification the generated data included one QTL up to six QTLs. This simulation study investigated 60 different QTL scenarios.

From the simulation study, the stochastic search identified only the QTL with no false positives for every simulation from 1 to 6 QTL and all effect sizes. Table I illustrates the effectiveness of the proposed approach to find the marker across multiple chromosomes without getting stuck in local troughs of nearby markers.

TABLE I  
SUMMARY RESULTS

Ground Truth	Effect Size	Gamma Noise ( $\alpha, \beta$ )	Proposed Results
C1M2	1	0.5,1	C1M2
		1,3	C1M2
C1M2	9	0.5,1	C1M2
		1,3	C1M2
C1M5, C2M15	2,4	0.5,1	C1M5, C2M15
		1,3	C1M5, C2M15
C1M5, C2M15	6,8	0.5,1	C1M5, C2M15
		1,3	C1M5, C2M15
C1M6, C2M15, C3M21	2,4,8	0.5,1	C1M6, C2M15, C3M21
		1,3	C1M6, C2M15, C3M21
C1M2, C1M9, C2M15, C5M31	1,3,5,9	0.5,1	C1M2, C1M9, C2M15, C5M31
		1,3	C1M2, C1M9, C2M15, C5M31
C1M2, C1M5, C1M9, C2M15, C4M27, C5M33	1,2,5,7,8,9	0.5,1	C1M2, C1M5, C1M9, C2M15, C4M27, C5M33
		1,3	C1M2, C1M5, C1M9, C2M15, C4M27, C5M33

A full disclosure of all experimental results is not possible due to the number of experiments conducted and the page constraints. However the interested reader may contact the lead author for the detailed results.

#### IV. CONCLUSION

The identification of QTL leads to discoveries that can impact society, e.g. identifying drought resistance for a plant or mass marker for cattle that can be used to breed meatier cattle over the use of steroids or bio-feed. Data obtained through QTL experiments are complex and methodologies need to be able to handle the complexities in a robust manner; hence, this research presented an approach based on Bayesian hierarchical model that tolerates complexity well. The Bayesian hierarchical model has shown that it is a flexible model that can incorporate multi-levels of information, e.g. trait values for recombinant plant lines or environmental information or laboratory information, in an experiment. The proposed approach was able to *correctly identify every* QTL for each of the 60 experiments conducted. The algorithm appears to be robust to the assumption of normality, although further investigations might be needed.

Future work will investigate the use of this method to model epistasis among markers and environmental factors.

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