Graph theoretical approach to study eQTL: a case study of Plasmodium falciparum

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ABSTRACT

Motivation: Analysis of expression quantitative trait loci (eQTL) significantly contributes to the determination of gene regulation programs. However, the discovery and analysis of associations of gene expression levels and their underlying sequence polymorphisms continue to pose many challenges. Methods are limited in their ability to illuminate the full structure of the eQTL data. Most rely on an exhaustive, genome scale search that considers all possible locus-gene pairs and tests the linkage between each locus and gene.

Result: To analyze eQTLs in a more comprehensive and efficient way, we developed the Graph based eQTL Decomposition method (GeD) that allows us to model genotype and expression data using an eQTL association graph. Through graph-based heuristics, GeD identifies dense subgraphs in the eQTL association graph. By identifying eQTL association cliques that expose the hidden structure of genotype and expression data, GeD effectively filters out most locus-gene pairs that are unlikely to have significant linkage. We apply GeD on eQTL data from Plasmodium falciparum, the human malaria parasite, and show that GeD reveals the structure of the relationship between all loci and all genes on a whole genome level. Furthermore, GeD allows us to uncover additional eQTLs with lower FDR, providing an important complement to traditional eQTL analysis methods.

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1 INTRODUCTION

The development of methods that allow us to uncover mechanisms of gene regulation and reconstruct gene regulatory networks is an important open problem in molecular biology. The advancement of high-throughput genotyping and gene expression platforms supports the analysis of expression quantitative trait loci (eQTL) as a tool to elucidate gene regulation. eQTL analysis considers expression of each gene as a quantitative trait and maps it to a genomic locus or marker. The genotype associated with a gene’s expression level highlights the genome region carrying the DNA polymorphism impacting the expression. The polymorphism may reside in the gene’s coding region or in a transcription factor binding site and could affect the expression level of its own or other genes in an inheritable way (Brem and Kruglyak, 2005; Monks et al., 2004; Petretto et al., 2006). Hence, a significant statistical linkage between a locus and a gene’s expression suggests that the gene in question is regulated by the locus, which may hold a regulatory element or a regulator gene. Since the early work of Jansen and Nap (Jansen and Nap, 2001), eQTL has become a widespread technique to identify such regulatory associations and has been applied to several species including yeast (Brem and Kruglyak, 2005; Yvert et al., 2003), mouse (Bystrykh et al., 2005; Chesler et al., 2005) and human (Cheung et al., 2005; Stranger et al., 2005). Typically, these studies use genome-wide association studies (GWAS), considering loci spanning the genome and expression profiles of all genes in the organism. As a major advantage, simultaneous monitoring of thousands of gene expression traits provides unique and unbiased data and opens the possibility of constructing a global view of the underlying regulation machinery.

Despite the valuable insights that can be gained, current attempts to elucidate the structure of eQTLs still face many challenges. Only a few methods are available that model complete eQTL data to discover broader eQTL structure. The complex dependence of the variations of gene expression regulation on phenotypic differences nurtures the expectation that important information can be gained from considering more subtle relationships between genotype and expression. The large number of gene expression traits and genomic loci poses challenges for both computational efficiency and statistical power. Traditionally, an eQTL study tests the linkage between all genes’ expression and all loci, adding up to millions of single statistical tests. For example, (Stranger et al., 2007) used 2 million single-nucleotide polymorphism (SNP) and more than 13,000 transcripts, leading to more than 10^{10} tests for all possible associations, a number that causes a serious multiple testing issue (i.e. the chance of false positives in a family of multiple hypothesis tests is higher than that of a single test). Consequently, that study was restricted to consider mainly cis-regulation—associations between SNPs and genes within 1 Mb of the SNP in question—which reduced the number of tests dramatically. While more complex regulation programs are of increasing interest (Storey et al., 2005), the combinatorial nature of such problems and the large number of loci call for improved methods that allow discovery of more complex regulation programs involving more than one locus and one gene.

To address such problems, we propose a novel method, GeD (Graph based eQTL Decomposition), to analyze eQTL data. Our method models the genotype, progeny and expression data as an eQTL association graph, a three–partite graph which is the union of two bipartite graphs. By simultaneously exploring two bipartite graphs, GeD discovers sets of dense subgraphs, called

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eQTL association cliques, each containing a set of loci, a set of genes and a set of genotypes and a set of genes. The progenies provide evidence that the set of loci may be associated with the set of genes. Such eQTL association cliques give a succinct representation of structures among loci, progenies and genes on a genome-wide scale. More importantly, each locus, progeny and gene can appear in more than one association clique, which depicts a complete picture of eQTL data. To find eQTL association cliques, GeD employs an efficient bipartite clique enumeration algorithm initially designed for building a concept lattice (Farach-Colton and Huang, 2008). The set of association cliques helps to select a small set of locus–gene pairs that are expected to have significant linkage; subsequently, statistical tests, including corrections for multiple testing, are performed for these selected locus–gene pairs.

Testing all possible locus–gene pairs, and new eQTL hotspots 1327 eQTLs in different chromosomes tend to co-occur in some eQTL association significantly different from random association cliques, and loci on ≤ outlined here. Despite these challenges, Gonzales and increases the need for the more discerning methodology exist, although the association between loci and gene expression P. falciparum specific transcription factors in parasite is of fundamental importance, successes in identifying eQTL association cliques, each containing a set of loci, a set of genes, and a set of genotypes and a set of genes. The progenies provide evidence that the set of loci may be associated with the set of genes. Such eQTL association cliques give a succinct representation of structures among loci, progenies and genes on a genome-wide scale. More importantly, each locus, progeny and gene can appear in more than one association clique, which depicts a complete picture of eQTL data. To find eQTL association cliques, GeD employs an efficient bipartite clique enumeration algorithm initially designed for building a concept lattice (Farach-Colton and Huang, 2008). The set of association cliques helps to select a small set of locus–gene pairs that are expected to have significant linkage; subsequently, statistical tests, including corrections for multiple testing, are performed for these selected locus–gene pairs.

Fig. 1. (a) Each genomic locus I is assigned the genotype of the strain it was inherited from (0/1). In all strains s, we discretize expression levels of genes g as being either ‘up’, ‘unchanged’ or ‘down’. Considering the genotypes of I, we observe different gene expression patterns, indicating different expression mechanisms in different strains. For example, the expression of g1 is up-regulated in strain s1, s2, s3 and s6, and down-regulated in strain s2, s3, s4 and s6. In (b) we show the corresponding eQTL association graph. Specifically, we find an association clique including genes g3 and g5, strains s2, s3, s4 and s6, and loci I0 and I0, shown in light grey. The edges of the association clique are drawn with wider lines.

2 MATERIALS AND METHODS

First we introduce the basic rationale of the GeD approach and present a detailed description of GeD. Finally, we describe the P. falciparum eQTL data we used to test our method.

2.1 Problem definition

In an eQTL experiment, we consider a set of progeny strains, often obtained from a cross between two parental strains with different genetic and phenotypic background. In our case, we only consider two possible genotypes (0, 1) for each locus and assign each locus l to the parental strain the locus was inherited from. All strains can be partitioned into two groups by the genotype of a given locus, and we discretize the expression levels of each gene as being either ‘up-regulated’, ‘unchanged’ or ‘down-regulated’ (Fig. 1a).

To represent the above relationship between loci and genes, we define an eQTL association graph G(Gs,Us,Le,Ep) as follows: The graph contains three sets of vertices (Gs, Us, Le), where Gs represents the genotypes of loci, Us represents progeny strains, and Le represents up- or down-regulated gene expression. Vertices g and u indicate a gene g’s up- or down-regulation and u and l represent the genotype at locus l as either 0 or 1. An edge between g∈G and u∈U and a progeny strain s indicates g’s expression is up- or down-regulated in strain s. An edge between lj∈L and a progeny strain s indicates the genotype of lj is 0 in strain s. Note, that there are no edges between genes G and loci L. Our eQTL association graph can be viewed as a three-partite graph which is the union of two bipartite graphs BG1(L,U,s1) and BG2(G,U,s2).

The corresponding eQTL association graph in Figure 1a is shown in Figure 1b, where we consider the subgraph induced by g1, g2, g3, g4, g5, g6, s1, s2, s3, s4, s5, s6, lj0, lj1 and I0. We call such a subgraph an eQTL association-clique, defined as To = (G1∪S1∪L1,E1), where g1∈G1, u1∈U1, lj0∈L1, and ∀(g1,u1)∈E1. In other words, we require that G1, S1, and L1 are fully connected. Additionally, no such association clique To = (G2∪S2∪L2,E2) exists, where G2 and S2 are fully connected, L2 and S2 are fully connected, and G2∪S2∪L2 is a maximal subgraph that cannot be extended further, maintaining full connectivity. Similarly, an eQTL association clique can be viewed as the union of two dense bipartite subgraphs formed by Gp∪Sp, and Lp∪Sp, respectively. As defined, please note that in each eQTL association clique, |G| ≥ 1 and |G| ≥ 1. Furthermore, opposing loci I0 and I1, or gene expression states g0 and g1 cannot appear in the same association clique.

It is easy to see that there can be four cases where a locus–gene pair (lj, g) can appear in an association clique: The first case is that an up-regulated gene g0 and a 0-genotyped locus lj0 are in an association clique while in the second case a down-regulated gene g1 and a 1-genotyped locus lj1 are in an association clique. We call these cases P1. In a third case an up-regulated gene g0 and a 1-genotyped locus lj0 are in an association clique while in the last case a down-regulated gene g1 and a 0-genotyped locus lj1 are in an association clique, cases we call P2. We call the first two cases compatible since they both suggest that g’s expression pattern is different in two groups defined by g’s genotype—and vice versa.

Intuitively, a locus and a gene that co-appear in an association clique that has a large subset of strains are expected to be more closely associated. Therefore, we define the size of the progeny strain set in a subgraph of the association graph as support, sp. For a locus–gene pair (lj, g) and an association clique with support sp, if gj and lj co-appear in the clique, we define the support provided by the clique sp. Similarly, we define
Analogously, the support for $P_2$ is $sp_2 = \max(p_{21}, p_{22}) + \max(p_{23}, p_{24})$ over all eQTL association cliques. Analogously, the support for $P_1$ is $sp_1 = \max(p_{11}, p_{12}) + \max(p_{13}, p_{14})$. Since $P_1$ and $P_2$ are opposites if we consider the linkage between $g_i$ and $g_j$, we define $sp = sp_1 - sp_2$ as a rough measurement of the net support for the expectation that significant linkage between $g_i$ and $g_j$ exists.

2.2 Method

Based on these important heuristics, GeD performs the following steps to identify eQTL association cliques and to detect eQTL:

(i) Discretize (see below) gene expression levels and build an eQTL association graph $G_{\text{E}}(V(G_{\text{E}}), E_{\text{E}})$, a union of bipartite graphs $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$ and $B_{\text{G}}(V(B_{\text{G}}), E_{\text{E}})$. 

(ii) Find all maximal bipartite cliques in $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$ for each sample set $S_i$ in the bipartite graph induced by $S_i \cap B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$. 

(iii) Identify sets $G_i$, where each vertex is connected to each vertex in $E_i$ appearing in $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$. If the subgraph $\text{E}_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$ has not been generated yet, output this graph as an eQTL association clique.

(iv) For each locus–gene pair $(l_i, g_j)$ appearing in one eQTL association clique, select the pair if its support value $\max(p_{ij1}, p_{ij2}) + sp$ meet criteria described below.

(vi) Among selected locus–gene pairs compute $p$-values of their association (adjusted for multiple testing).

In both steps (ii) and (iii), it is essentially to enumerate bipartite cliques from a large bipartite graph efficiently. We apply an algorithm for building a concept lattice, which can be considered a hierarchical structure for organizing all bipartite cliques given a bipartite graph. Such structures have been used to compute gene expression matrices (Huang and Parach-Cohn, 2007). The delay-time complexity—the time spent to compute each bipartite clique—of the algorithm is $O(n^2)$, where $n_1$ and $n_2$ are the size of two sets of vertices in the bipartite graph.

Here, we assume that the number of bipartite cliques in $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$ is less than in $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$. If this is not the case, GeD starts from $B_{\text{E}}(V(B_{\text{E}}), E_{\text{E}})$ in step (ii); steps (iii) and (iv) are changed accordingly.

To obtain corrected $p$-value in the last step of GeD, we apply the method of Churchill and Doerge (Churchill and Doerge, 1994). For each gene $g_j$ in a selected locus–gene pair from step (v), we maintain a locus list $(l_{i1}, l_{i2}, ..., l_{in})$, where each locus in the list appears with $g_j$ in one of selected locus–gene pairs. We randomly permute $g_j$’s expression and compute the nominal $p$-value for the linkage between the random expression and a locus in the list and retain the smallest $p$-value. After repeating the process 1000 times, we use all retained $p$-values to approximate a null distribution. By comparing the nominal $p$-value from real data to the null distribution, we obtain the corrected $p$-value.

While in many ways to discretize gene expression data (Bricquet et al., 2002) transcription patterns of most genes in several major P-falciparum strains are very similar (Llinas et al., 2006). Therefore, we used a simple method (Quackenbush, 2002) that can be readily applied to our case. We computed the mean $\bar{m}$ and standard deviation $sdev$ for each probe and define genes with expression levels $> m + 3sdev$ as ‘up-regulated’ and $< m - 3sdev$ as ‘down-regulated’. Specifically, we set $b$ to 1, allowing us to detect more variation in the gene expression. Another advantage of $b=1$ is that each probe will be represented by at least one vertex in the association graph. In the worst case, the number of bipartite cliques in a bipartite graph is $\min(2^n, 2^m - 2)$, where $n_1$ and $n_2$ are the sizes of the two vertex sets of the bipartite graphs. Since thousands of vertices in $G$ and $E$ in the eQTL association graph exist, extreme computational costs we only allow bipartite cliques with at least five progeny strains in step (ii) and (iii).

2.3 Materials

Utilizing P-falciparum eQTL data from the reference (Gonzales et al., 2008), we used 34 progeny strains obtained from a HB3xDd2 cross. Each progeny was genotyped at 329 microsatellite markers along 14 chromosomes. Expression levels were measured 18 h after the parasite invades human erythrocytes (RBCs), by 7665 probes representing 5150 ORFs.

3 RESULTS

As previously mentioned, eQTL association cliques allow us to determine the structure inherent in eQTL data. We first show the difference between association cliques obtained from the underlying eQTL data as well as from randomized data. Subsequently, we report eQTLs we determine in eQTL association cliques.

3.1 Size Distribution of eQTL Association Cliques

Applying GeD, we obtain 135 044 eQTL association cliques with support $sp \geq 5$. Overall, the support in eQTL association cliques ranges from 5 to 10. To generate random eQTL association cliques, we permuted the expression vector of each probe and applied GeD with the same parameters on the random data 100 times. We find 40 773, 20 393 and 5396 association cliques with support of 5, 6 and 7 in the real data. In random data, we find on average more association cliques ($sp=5$: 77 200; $sp=6$: 28 019; $sp=7$: 5809). Applying a one-sample t-test between the number of association cliques in real and random data yielded $p < 10^{-11}$ in all three cases. Considering association cliques with $sp=8$, 9 and 10, we find 872, 84 and 5 in the real data. Compared to the random data, we analogously find significant differences. Specifically, we find on average 807, 74 and 4 random association cliques with the same supports in the randomized data with $p < 10^{-11}$ in the cases of support 8, 9 and $p < 10^{-10}$ in the case of support 10.

Subsequently, we compared the number of association cliques in real and random data that have the same support $sp$ and $|G|$, the number of probes. The number of association random association cliques was significantly smaller except when $sp=5$ or 6 or 7, $|G|=1$, and $sp=5$, $|G|=2$. In Figure 2a we show the number of real association cliques and the average number of random association eQTL with $sp=6$ and several different number of probes $|G|$. Specifically, the largest eQTL association clique with $sp=6$ has 87 probes.

A closer look revealed that, given support $sp$, $|G|$ and $|L|$, the number of association cliques in the random data was significantly larger than in the real data only when $|G|$ is small. For example, when $sp=6$ and $|L|=7$, the number of association cliques in the random data was larger only if $|G|=1$. For a given support value $sp$ and the number of loci $|L|$, the number of association cliques in the random data was significantly larger than in the real data for most cases when $sp=5$ or 6. We find similar results when $sp=7$, $|L| < 12$, $sp=8$, $|L| < 7$, and $sp=9$, $|L| < 5$. Specifically, we show the number of random association cliques with $sp=6$ and different numbers of loci $|L|$ in Figure 2b.

Since genotypes of adjacent loci in the same eQTL association clique are adjacent. Though this is frequently the case, we also find many co-appearing loci although they are on different chromosomes. For example, loci 2_0 on chromosome 2 and 12_45.8 on chromosome 12 co-appear with six loci on chromosome 3 in an eQTL association clique with support 10. We did not find such a result in the random data, suggesting that these loci tend to co-segregate and indicating
Fig. 2. Number of association cliques from real and random data with support 6. Numbers we obtained from randomized data were averaged over 100 runs. In (a) we perform the analysis varying |G|, the number of probes, while we show the analogous results with changing number of loci |L| in (b).

that a closer examination of their relation might be interesting with linkage disequilibrium based methods for P. falciparum (Su and Wootton, 2004).

3.2 eQTL detection

We have shown that a locus–gene pair appearing in two eQTL association cliques in a compatible way is more likely to have a significant linkage than those pairs that do not. Hence, we could use eQTL association cliques to select a small number of locus–gene pairs to be tested for linkage, many of which we expect to yield significant $p$-value. To select locus–gene pairs $(l_j, g)$, we use criteria $\max(s_{P_1}^{l_j}, s_{P_2}^{l_j}) \geq 12$ and $s_{P_j} \geq 6$ in step (v) to select locus–gene pairs $(l_j, g)$. Please note that each association clique has at least five progeny strains because we generate maximal bipartite cliques with at least five progeny strains. If we assume a locus–gene pair $(l_j, g)$ with pattern $P_1$, then the minimum value for $s_{P_2}^{l_j}$ is 10 since $s_{P_2}^{l_j} = \max(s_{P_1}^{l_j}) + \max(s_{P_0}^{l_j})$, where $s_{P_0}^{l_j} \geq 5$ and $s_{P_0}^{l_j} \geq 5$. Note, that if we set this threshold too high, we potentially remove locus–gene pairs having significant linkage. In total, we selected 6232 locus–gene pairs. Figure 3 shows the histogram of nominal $p$-value computed by a two-sided $T$-test for the linkage of these selected pairs and all possible locus–gene pairs.

We observe that selected locus–gene pairs from association cliques yield significantly lower linkage $p$-value. Correcting $p$-values (see Methods section) and calculating FDRs (Storey and Tibshirani, 2003) we identified 2853 eQTLs ($p < 0.05$, FDR $< 0.04$). Identifying the most significant eQTLs, we use our set of association cliques we found in randomized data. With the same criteria, we obtained 6232 locus–gene pairs from each set of randomized association cliques and applied a $T$-test to obtain $p$-values for these pairs. In this way, we obtained 100 groups of $p$-values from random data, allowing us to estimate an empirical null distribution, which is often more stringent than the null distribution obtained individually for each gene (Churchill and Doerge 1994).

We required that each reported eQTL has a nominal $p$-value smaller than 90% of $p$-values in the empirical null distribution. Following this protocol, we found 1327 eQTLs for 513 probes (482 genes) and 231 loci. Previously, Gonzales et al. (Gonzalez et al., 2008) identified a set of 1063 eQTLs with FDR $< 0.24$ using standard GWAS. Figure 4 shows the distributions of eQTLs identified by Gonzales et al. and 1327 eQTLs we obtained with GeD. We observe that the distribution of eQTLs detected by GeD is similar to the distribution of previously identified eQTLs, which were obtained by considering all possible locus–gene pairs. We also find 251 (≈25%) eQTLs that appear in both sets. Although the overlap is considerable the two sets are quite different, an observation that can be attributed to fundamental differences in the methods (see Discussion section). Both analyses show that there are several eQTL hotspots on chromosome 3, 5 and 7. Gonzales et al. (Gonzalez et al., 2008) called a locus eQTL hotspot if there existed at least 14 linked probes at a particular locus. Analogously, we found 17 eQTL hotspots and discovered two/three new eQTL hotspots in the right/left subtelomeric region on chromosome 3. While two weak eQTL hotspots on chromosome 9 and 12 detected by Gonzales et al. did not appear in our result, we detected two new eQTL hotspots on chromosome 5 and 7. Note that the definition of a hotspots used in both studies does not differentiate between cis- and trans- links, and...
the reported hotspots represent the combined effect of both types of regulation as well as that of the pattern of linkage disequilibrium.

Both subtelomeric regions on chromosome 3 are enriched with highly polymorphic surface antigen genes such as cytoadherence linked asexual genes (CLAG), stevor genes, and var genes (Gardner et al., 2002). While compelling, it remains to be experimentally determined if such polymorphic antigen genes are indeed regulated by eQTL hotspots we identified in the same region. Interestingly, it has been reported that the right telomere of chromosome 3 has an extended region of similarity with the right telomere of chromosome 2, and some pseudogene sequences in the regions were also preserved (Bowman et al., 1999). Such preservation in these rapidly evolving regions may imply that these subtelomeric regions are biologically significant (Bowman et al., 1999), suggesting that the detection of additional eQTL hotspots in these regions provided more evidence for their importance in regulating the host-parasite interface. We also performed Gene Ontology term enrichment analysis for the target genes of newly detected eQTL hotspots using GOTermFinder (Boyle et al., 2004). We found that two hotspots show enriched GO terms referring to drug interaction and parasite-human invasion. The GO annotation of target genes of eQTL at locus 5p 25.8 on chromosome 5 was enriched for cytoadherence to microvasculature mediated by parasite protein and interaction with the host (p < 0.03).

4 DISCUSSION

We introduced a novel method—GeD—that integrates genotype, expression and progeny data, providing an analytical framework for the determination of gene regulation programs. In an eQTL association clique, vertices representing a locus’ genotype are fully connected with vertices that represent progeny strains. Such a structure refers to the case that loci have the same genotype when restricted to these progeny strains. Analogously, vertices that represent genes are fully connected with vertices representing progeny strains, indicating that the corresponding progeny strains share the same gene expression patterns. As such, eQTL association cliques allow the determination of associations of loci, progeny strains and genes in a simple way. In addition, the number of progeny strains supports the linkage between loci and genes in the same association clique, which can help to detect eQTLs.

In this article we focused on the application of the eQTL association cliques to enhance eQTL discovery. However, eQTL association cliques have the potential to answer other questions as well. For example, loci that are not in linkage disequilibrium and co-occur in a highly supported clique might indicate functionally important co-segregation. Note that while loci that are in the same clique and are genomic neighbors are likely to be in linkage disequilibrium. However, the opposite case is not necessarily true. This observation should be useful in elucidating non-random properties of linkage disequilibrium. Additionally, eQTL association cliques may help the identification of loci and genes that are related in a certain phenotype. If the phenotype of progeny strains in an association clique is different from remaining progeny strains, the loci and genes in the corresponding association clique are the prime candidates that affect the phenotype in question.

Using eQTL association cliques might also help to uncover multiple locus linkage. For example, consider loci l, and l' and gene g, and four eQTL association cliques, where l0 and l1 appear with g0, in one clique, l0 and l1 appear with g0 in another clique, l0 and l1 appear with g0 in the third clique and l0 and l1 appear with g0 in the last clique. It is unlikely that l0 or l1 are associated with g0 individually because the genotype 01 of l0 is associated with both up- and down-regulated expression of g0. The same rational holds for locus l. But since the joint genotype 01 of l0 and l1 is associated with up-regulation of g0’s expression, and joint genotype 01 and 10 of l0 and l1 is associated with down-regulation of g0’s expression, the two loci can have a significant epistatic interaction effect on g0. By restricting our attention on loci in the same association clique, we can select a small set of triplets (l0, l1, g0), which fit the above scenario, by simply counting association cliques. Testing the selected triplets for epistatic effects reduces the number of statistical tests, \( O(L^3|G) \), required by an exhaustive search, where \( L \) is the locus set and \( G \) is the gene set.

In our method, we modeled underlying data using certain choices. First, discretizing expression data, a gene was considered differentially regulated if its expression level was at least one standard deviation away from its mean expression. This choice was dictated by its relative simplicity and applicability of that method to the data where differences in the expression levels are not expected.
to be very large. Other methods of discretizing expression data would be considered in the future improvement of the method. Next, we chose to look at maximal cliques rather than other densely connected subgraphs, allowing us to avoid the introduction of additional ‘density’ parameter. Furthermore, such an approach also allowed us to easily generate such clique-structures utilizing the efficient bipartite clique enumeration method (Farach-Colton and Huang, 2008). While bipartite cliques can potentially be replaced with bi-clusters, the best heuristic for the identification of such overlapping bi-clusters remains to be found. We conclude that our choices might potentially influence our ability to detect potential eQTLs. However, we made our choices as simple as possible and highlight the usability of our novel method.

We applied GeD to progeny data of *P.falciparum* and found that eQTL association cliques have very different structures and distributions compared to random association cliques. Using eQTL association cliques to select a small set of locus–gene pairs, we corroborated previously identified eQTLs, and significantly increased their number, including new eQTL hotspots. Preliminary analysis of the possible functional relevance of these new eQTL hotspots showed that some harbor important antigen genes while others include target genes involved in drug and parasite-host interactions. Compared to previous results, we conclude that GeD bolsters traditional eQTL analysis methods and provides new opportunities for the discovery of critical biological functions in *P.falciparum*. Approximately 25% of eQTLs in the two eQTL sets identified by GeD and Gonzales et al. (Gonzales et al., 2008) overlap, a difference that can be caused by several factors. First, Gonzales et al. applied an interval mapping method based on a complex Bayesian model for QTL detection (Sen and Churchill, 2001). Assuming each marker is the potential eQTL location, we will allow us to tackle more challenging GWAS problems. Furthermore, some in human studies where we have larger eQTL data sets containing thousands or even millions of loci and genes in the worst case. Therefore, the scalability of GeD to association cliques can increase exponentially with the number of loci and genes in the worst case. Therefore, the scalability of GeD to association cliques can increase exponentially with the number of association cliques. In contrast, the GWAS used by Gonzales et al. is likely to miss more subtle associations detected by our method because only the most significant eQTLs can pass multiple testing correction performed for all possible locus–gene pairs.

Our current implementation of GeD is designed for the analysis of the large data set of *P.falciparum*. However, the number of eQTL association cliques can increase exponentially with the number of loci and genes in the worst case. Therefore, the scalability of GeD to larger eQTL data sets containing thousands or even millions of loci remains to be tested. Specifically, in human studies, where we have to deal with huge amount of expression and genomic data we expect strongly increasing computational costs, prompting the development of further heuristics and improved computational techniques that will allow us to tackle more challenging GWAS problems.

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