Chromosome mapping: radiation hybrid data and stochastic spin models

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Abstract

This work approaches human chromosome mapping by developing algorithms for ordering markers associated with radiation hybrid data. Motivated by recent work of Boehnke et al. [1], we formulate the ordering problem by developing stochastic spin models to search for minimum-break marker configurations. As a particular application, the methods developed are applied to 14 human chromosome-21 markers tested by Cox et al. [2]. The methods generate configurations consistent with the best found by others. Additionally, we find that the set of low-lying configurations is described by a Markov-like ordering probability distribution. The distribution displays cluster correlations reflecting closely linked loci.
I. INTRODUCTION

The use of data from radiation hybrid (RH) experiments has become a useful method for fine structure mapping of human chromosomes. Based on methods described by Goss and Harris [3, 4], Cox et al. [2] and Burmeister et al. [5] have developed the technique in detail so that results from their experiments provide material for ordering DNA markers on human chromosomes.

The basic strategy employed in radiation hybrid mapping (fully described in Cox et al. [2]) entails irradiating a rodent-human somatic cell hybrid, that contains a particular human chromosome, with a lethal dose of x-rays. This will cause the chromosomes to break into several fragments. After fusion with HPRT-deficient rodent cell lines, only the fused cells, containing both the X-irradiated cells and the normal rodent cells, will survive if grown in HAT medium. Each hybrid clone arising from this fusion will contain a unique set of fragments from the original human chromosome. Each clone can then be typed for a set of human DNA markers (equivalently, loci) known to be on that human chromosome. Based on the assumption that tightly linked markers are unlikely to be broken apart by the radiation, markers close to one another will show a correlated pattern of retention in the clones; whereas, distant markers will be retained in a relatively independent manner.

Several methods for ordering markers have been developed using results from RH experiments [including both parametric (Cox et al. [2]; Bohnke et al. [1]) and non-parametric methods (Bohnke et al. [1]; Falk [6]; Bishop and Crockford [7]; Weeks et al. [8]). In particular, Bohnke et al. [1] used a mathematical quantity associated with the number of breaks and then used optimization techniques to minimize that quantity. One optimization
technique involved a simulated annealing search for configurations associated with minimal numbers of breaks.

We set out to study and understand the work of Boehnke et al., and we developed a formulation in the context of stochastic spin systems (Falk [9]). It may be useful to point out some differences in the formulations.

Boehnke et al. use block inversions of a given marker order and compare the old and new orders with respect to "obligate" breaks. They then apply simulated annealing techniques and decide, at each time step, whether to retain the original order or transition to the new. If a transition would result in a smaller number of breaks, the transition is made with probability one. If a transition would not decrease the number of breaks, then the transition probability is less than one, and that transition probability systematically decreases over time.

In our study we implement three algorithms which incorporate the number of "breaks". The three algorithms are three stochastic spin models. These, too, are designed to search for configurations with small numbers of breaks. A probability is constructed to determine whether or not to retain the current order or transition to the new. The probabilities are set up so as to bias the decision towards transitions to configurations with fewer breaks; however, at a given step, a possible transition leading to a smaller number of breaks, will not necessarily be realized.

The spin language provides mathematically intuitive expressions for the number of breaks. Those expressions contain products of adjacent spin variables, and calculations involving breaks are easily presented in spin notation.
For those seeking a rigorous mathematical setting, we remark that Liggett [10] has treated stochastic spin models and related models from biology, physics and econonomics. Liggett’s book contains an extensive bibliography and guides the reader to survey papers by Griffeath, Durrett, Stroock, Holley, and others.

II. METHODS

We are considering $M$ clones, each of which is tested for the presence (or absence) of $N$ different DNA markers. It is convenient to represent the clones as $M$ rows, each with $N$ sites. Thus, one pictures a two-dimensional array of $M$ rows, $N$ columns.

\[
\begin{array}{cccc}
1 & 2 & \cdots & N \\
1 & \bullet & \bullet & \bullet & \cdots & \bullet & \bullet \\
2 & \bullet & \bullet & \bullet & \cdots & \bullet & \bullet \\
\vdots & \vdots & \ddots & \ddots & \ddots & \ddots & \ddots \\
M & \bullet & \bullet & \bullet & \cdots & \bullet & \bullet \\
\end{array}
\]

Assign a particular DNA marker, labelled $f_j$, ($j = 1, 2, \ldots, N$), to each column. Associate a variable ("a spin") $s_{ij}$ with the site at row $i$, column $j$. If the marker $f_j$ is present at site $(i, j)$, take $s_{ij} = +1$; if $f_j$ is not present at site $(i, j)$, take $s_{ij} = -1$. If one is uncertain as to whether a marker is present at site $(i, j)$, we take $s_{ij}$ to be unknown, and we deal with such sites in a manner to be specified subsequently.

For ($j = 1, 2, \ldots, N$) the marker $f_j$ was arbitrarily assigned to column $j$. But any of the $N!$ assignments would be possible. A criterion is needed for
judging the "goodness" of an assignment.

Considering that the DNA markers being tested for lie on a particular human chromosome, those markers which are tightly linked are likely to appear together or not appear in each clone. Therefore, it is reasonable to seek those marker assignments which reflect such correlation. For that purpose we say that a "break" exists between sites \((i, j)\) and \((i, j + 1)\) whenever \(s_{ij}s_{i,j+1} = -1\). The strategy is to minimize the total number of breaks. Note that in the two-dimensional array of spins, a break refers only to horizontal, nearest-neighbor spin pairs.

Here we devise and test several algorithms which monitor the total number of breaks while selectively permuting column labels. The algorithms attempt to explore the vast configuration space in the manner of a "random walk", biased toward configurations having a reduced number of breaks. Notice, for \(N = 14\) there are \(N! = 87178291200\) permutations of the numbers \((1, 2, ..., 14)\). Thus, in the spirit of the travelling salesman problem, simulated annealing techniques are used (Kirkpatrick et al [11], Press, et al [12]).

**Nearest-neighbor transposition algorithm**

(0) Start with a specified configuration \(\{s\}\) of the variables \(s_{ij}\).

(1) Select a number \(j\) at random from the set \(\{1, 2, ..., N\}\).

(2) If \(j \neq N\), compute the total number of breaks between columns \(j - 1, j\) and between columns \(j + 1, j + 2\). Denote that number by \(B_j(1)\):
\[ B_j(1) = \sum_{q=1}^{M} \left[ (1 - \delta_{j,1}) \frac{1 - s_{q,j-1}s_{q,j}}{2} + (1 - \delta_{j,N-1}) \frac{1 - s_{q,j+1}s_{q,j+2}}{2} \right] \]  

for \( j \in \{1, 2, ..., N-1\} \). The Kronecker delta contained in \( (1 - \delta_{i,j}) \) is inserted to handle "end effects" since column 1 has no left neighbor and \( N \) has no right neighbor.

3) Repeat (2) with the spin values \( s_{ij} \) and \( s_{i,j+1} \) interchanged for \( i = 1, 2, ..., M \), and denote the resulting sum by \( B_j(2) \) instead of \( B_j(1) \).

4) Then compute \( B_j \), the change in the number of breaks resulting from the interchange of the columns of spin values \( s_{ij} \) and \( s_{i,j+1} \).

\[ B_j = B_j(2) - B_j(1) \quad \text{for} \quad j \in \{1, 2, ..., N-1\}. \]  

(2.2)

5) Interchange DNA marker column assignments and the associated spin values for columns \( j \) and \( j + 1 \) with probability.

\[ w_{1j} = \frac{\exp \left( -\beta B_j/M \right)}{2 \cosh \left( \beta B_j/M \right)} \quad \text{for} \quad j \in \{1, 2, ..., N-1\} \]

\[ = 0 \quad \text{for} \quad j = N. \]  

(2.3)

Do not interchange DNA marker column assignments and the associated spin
values for columns $j$ and $j + 1$ with probability

$$w_{2j} = 1 - w_{1j}. \quad (2.4)$$

Note:

$$w_{2j} = \frac{\exp (+\beta B_j/M)}{2 \cosh (\beta B_j/M)} \quad \text{for } j \in \{1, 2, \ldots, N - 1\}$$

$$= 1 \quad \text{for } j = N. \quad (2.5)$$

The "inverse temperature" parameter $\beta \ (0 \leq \beta < \infty)$ is allowed to increase in an empirically determined manner as the algorithm is implemented. Clearly very large $\beta$ strongly favors transitions reducing the number of breaks; whereas, a small, positive value of $\beta$ makes transitions to increase or to decrease the number of breaks almost equally likely. Why not just take $\beta$ large at the outset? The answer is (Kirkpatrick et al [11], Press et al [12]) that setting $\beta$ large early in the calculation may cause the algorithm to get "trapped" in a local minimum before performing a significant search of configuration space.

Thus, the initially chosen value for $\beta$ and the manner of increasing that value constitute the "art" of simulated annealing.

(6) Return to (1) and repeat the procedure starting with the current configuration $\{s'\}$ of the variables $s'_{ij}$. The procedure may be terminated after
a specified number of iterations and/or when the total number of breaks

\[ B = \sum_{q=1}^{M} \sum_{j=1}^{N-1} \frac{1 - s_{qj}s_{q,j+1}}{2} \]  

(2.6)

has realized acceptably small values. For each run one retains the configurations associated with the smallest values of \( B \).

If \( \beta \) were fixed, then the above procedure would define a finite-state, discrete-time Markov chain with transition probability \( p(\{s'\} | \{s\}) \) from a configuration \( \{s\} \) to configuration \( \{s'\} \). Explicitly

\[
p(\{s'\} | \{s\}) = \frac{1}{N} \sum_{j=1}^{N} \left\{ \prod_{q=1}^{M} \left( \frac{1 + s'_{qj}s_{q,j+1}}{2} \right) \left( \frac{1 + s'_{q,j+1}s_{qj}}{2} \right) \right. \\
\times \left. \prod_{k=1; k \neq j, j+1}^{N} \left( \frac{1 + s'_{qk}s_{qk}}{2} \right) \right\} w_{1j} \\
+ \left. \left\{ \prod_{q=1}^{M} \prod_{k=1}^{N} \left( \frac{1 + s'_{qk}s_{qk}}{2} \right) \right\} w_{2j} \right\}. \]  

(2.7)

**Two-column transposition algorithm**

A natural extension of the nearest-neighbor transposition algorithm involves columns \( k, j \) with \( k > j + 1 \). Remove two numbers at random from the set \( \{1, 2, ..., N\} \). Denote the smaller number by \( j \) and the larger by \( k \). If
\( k = j + 1, \) follow the previously described nearest-neighbor algorithm, starting with step (2). If \( k > j + 1, \) proceed as follows:

Consider the total number of breaks between columns \( j - 1, j; j, j + 1; k - 1, k; k, k + 1. \) Denote that number by \( B_{jk}(1), \) where

\[
B_{jk}(1) = \sum_{q=1}^{M} \left[ (1 - \delta_{j,1}) \frac{1 - s_{q,j-1}s_{q,j}}{2} + \frac{1 - s_{q,j}s_{q,j+1}}{2} \\
+ \frac{1 - s_{q,k-1}s_{q,k}}{2} + (1 - \delta_{k,N}) \frac{1 - s_{q,k}s_{q,k+1}}{2} \right].
\tag{2.8}
\]

After interchanging spin values \( s_{ij} \) and \( s_{ik} \) for \( i = 1, 2, \ldots, M, \) the number of breaks is

\[
B_{jk}(2) = \sum_{q=1}^{M} \left[ (1 - \delta_{j,1}) \frac{1 - s_{q,j-1}s_{q,k}}{2} + \frac{1 - s_{q,k}s_{q,j+1}}{2} \\
+ \frac{1 - s_{q,k-1}s_{q,j}}{2} + (1 - \delta_{k,N}) \frac{1 - s_{q,j}s_{q,k+1}}{2} \right].
\tag{2.9}
\]

Then the change (in the number of breaks) resulting from the interchange is denoted by \( B_{jk}, \) where

\[
B_{jk} = B_{jk}(2) - B_{jk}(1).
\tag{2.10}
\]
Now interchange DNA marker column assignments and the associated spin values for columns \( j, k \) for \( k > j + 1 \) with probability

\[
w_{1jk} = \frac{\exp\left(-\beta B_{jk}/M\right)}{2 \cosh\left(\beta B_{jk}/M\right)} \quad \text{for } j \in \{1, 2, ..., N - 2\}.
\] (2.11)

Do not interchange DNA marker column assignments and the associated spin values for columns \( j \) and \( k \) with probability

\[
w_{2jk} = 1 - w_{1jk}.
\] (2.12)

**Block-flip algorithm**

As in the preceding algorithm, remove two numbers at random from the set \( \{1, 2, ..., N\} \). Denote the smaller number by \( j \) and the larger by \( k \). If \( k = j + 1 \), follow the previously described nearest-neighbor algorithm, starting with step (2). If \( k > j + 1 \), proceed as follows:
The block consists of columns $j, \ldots, k$. Before flipping the block, the number of breaks involving columns $j - 1, j$ and columns $k, k + 1$ is

$$B^{\text{block}}_{jk}(1) = \sum_{q=1}^{M} \left[ (1 - \delta_{j,1}) \frac{1 - s_{q,j-1}s_{q,j}}{2} + (1 - \delta_{k,N}) \frac{1 - s_{q,k}s_{q,k+1}}{2} \right].$$

(2.13)

After flipping the block, the number of breaks is

$$B^{\text{block}}_{jk}(2) = \sum_{q=1}^{M} \left[ (1 - \delta_{j,1}) \frac{1 - s_{q,j-1}s_{q,k}}{2} + (1 - \delta_{k,N}) \frac{1 - s_{q,k}s_{q,k+1}}{2} \right].$$

(2.14)

Then the change in the number of breaks is denoted by $B^{\text{block}}_{jk}$, where

$$B^{\text{block}}_{jk} = B^{\text{block}}_{jk}(2) - B^{\text{block}}_{jk}(1).$$

(2.15)

Now flip the block with probability
\[ w_{1jk}^{\text{block}} = \frac{\exp(-\beta B_{jk}^{\text{block}}/M)}{2 \cosh(\beta B_{jk}^{\text{block}}/M)} \quad \text{for } j \in \{1, 2, \ldots, N-2\}. \quad (2.16) \]

Do not flip the block with probability

\[ w_{2jk}^{\text{block}} = 1 - w_{1jk}^{\text{block}}. \quad (2.17) \]

The preceding method is similar to the block inversion algorithm used by Boehnke et al., but our transition probabilities differ from theirs.

**Unknown site content**

In any clone one may have strings of one or more sites where the DNA marker associated with each site in a string is unknown. Thus, the spin values are unknown for the string. In the above algorithms such unknowns are dealt with in the following way.

Consider the case where the left and right ends of a string connect respectively to known spin \( s_{\text{left}} \) and to known spin \( s_{\text{right}} \). If \( s_{\text{left}} = s_{\text{right}} \), then all spins in the string are replaced by \( s_{\text{right}} \). If \( s_{\text{left}} \neq s_{\text{right}} \), then a fair coin toss is simulated. If the coin shows head (tail), then all spins in the string are replaced by +1 (−1).
If a string (of unknowns) contains an end spin, then replace all spins in
the string by the value of the connecting spin \(s_{\text{left}}\) or \(s_{\text{right}}\), as appropriate.

With all spin values now specified, the number of breaks can be calculated
for any of the above algorithms, and the relevant transition probability can
be evaluated. The simulated annealing proceeds one step. After that step,
all of the previously unknown spin values are again regarded as unknown.
[Note: Due to a possible column interchange or block flip, those unknown
spins, which previously belonged to particular strings, may now belong to
different strings.] The above prescription for replacing unknown spin values
by +1 or −1 is now repeated, and the transition probabilities are reevaluated.
The simulated annealing proceeds another step, etc.

**III. APPLICATION**

As an example, consider the data presented by Cox et al. [2] relating
to 14 markers on chromosome 21 that were tested in 99 RH clones. The 14
markers are given in Table 1. These are the same data used by Boehnke et
al. [1] and presented in detail in their Table 1. For our algorithms an entry
of 1 in their table becomes +1, 0 becomes -1 and a "?" remains unknown
and is treated at each step as described above.

All three algorithms were applied to the data in the 99 \(\times\) 14 matrix for
200,000 iterations. Initial values of \(\beta\) and incremental steps for increasing
\(\beta\) were chosen. This produced a set of permutations with acceptably small
values of B, the total number of breaks for a particular configuration. For
each run, a ranked set of marker permutations with the smallest values of B
was retained.
Table 2 lists the 2 distinct "best" orders found by each algorithm in a representative run. As can be seen, the first algorithm, where two nearest-neighbor columns are transposed, does not result in permutations with values of B that are as low as those reached by algorithms 2 and 3. Although, in principle, the nearest-neighbor transposition should allow for visiting all possible permutations of the columns, in practice, such exploration is not efficiently accomplished here. The large configuration space, and the empirical nature of selecting $\beta$ to implement simulated annealing provide a computational challenge for the first algorithm. The other two algorithms display improved ability to achieve low B values with the chosen set of parameters. Additionally these algorithms reach the same optimal order as that attained by Boehnke et al. (see their Table 2), with the same number of breaks. The only difference is that we have retained all 14 markers; whereas they combined markers S12 and S111, since the latter markers were indistinguishable in the data matrix. Hence for each marker order in their Table 2 we would have 2 orders.
Although algorithms such as these do not assure that the marker order with the smallest number of breaks is the correct order, inspection of a set of low-lying states leads to some useful information about the stability of clusters of markers that retain their local ordering. For example, consider the set of unique permutations representing the 24 "best" orders obtained from a series of runs of the three algorithms (Table 3). We see, e.g., that S47 and SOD are nearest neighbors in all 24 permutations and appear as the last two markers in 16 permutations. Similarly, the triplet S46-S4-S52 is preserved in 21 permutations and of these, positions 2, 3 and 4 contain S46, S4 and S52 respectively 15 times. Based on observations such as these, we looked for a general ordering property associated with sets of low-lying configurations.

**Markovian-like property of low-lying configurations**

Let the DNA marker at site $j$ be denoted by $f_j$, where $f_j$ is a member of the set \{S16, S48, S46, S4, S52,...\}. A configuration of sites is denoted by the ordered N-tuple $(f_1, f_2, ..., f_N)$. We have used underlining to distinguish a DNA marker such as S11 from a spin variable such as $s_{11}$.

For any configuration of sites, one can define strings $(f_i, f_{i+1}, ... f_{i+m})$ with $i = 1, ..., N; 0 \leq m \leq N - i$.

Consider a collection $\mathcal{C}$ of distinct configurations. In the collection the probability of a string, $(f_i, f_{i+1}, ..., f_{i+m})$ is denoted by $P_{i,...,i+m}(f_i, f_{i+1}, ..., f_{i+m})$.

Now consider the Markovian-like approximation
\[ P_{i,...,i+m}(f_i, f_{i+1}, ..., f_{i+m}) \approx P_{i,i+1,i+2}(f_i \mid f_{i+1}, f_{i+2}) P_{i+1,i+2,i+3}(f_{i+1} \mid f_{i+2}, f_{i+3}) \ldots \]

\[ \ldots P_{i+m-2,i+m-1,i+m}(f_{i+m-2} \mid f_{i+m-1}, f_{i+m}) P_{i+m-1,i+m}(f_{i+m-1}, f_{i+m}) \quad (\star) \]

for \( i = 1, ..., N - 2 \); \( 2 \leq m \leq N - i \).

**Example 1.**

Consider the collection of 24 distinct low-lying configurations given in Table 3. Let (the number of markers) \( N = 14 \). Look at the cluster \((f_6 = F, f_7 = G, f_8 = H, f_9 = I, f_{10} = J)\) where \( F \) denotes the DNA marker \( S_{11} \), \( G \) denotes \( S_1 \), \( H \) denotes \( S_{18} \), \( I \) denotes \( S_8 \), and \( J \) denotes \( \text{APP} \). This marker assignment corresponds to the ordering of the first row of Table 3.

For the above collection we find the frequency

\[ P_{6,7,8,9,10}(F, G, H, I, J) = 7/24 \quad (3.1) \]

and we also find the frequencies
\[ P_{6,7,8}(F \mid G, H) = \frac{11}{11} \] (3.2)

\[ P_{7,8,9}(G \mid H, I) = \frac{7}{11} \] (3.3)

\[ P_{8,9,10}(H \mid I, J) = \frac{11}{11} \] (3.4)

\[ P_{9,10}(I, J) = \frac{11}{24}, \] (3.5)

so the approximation (⋆) with \( i = 6, m = 4 \) is satisfied by the observed frequencies.

Similarly, for the above configurations \( P_{7,8,9,10}(G, H, I, J) = \frac{7}{24} \) and

\[ P_{7,8,9,10}(G, H, I, J) \approx P_{7,8,9,10}(G \mid H, I)P_{8,9,10}(H \mid I, J)P_{9,10}(I, J) \]

\[ = (\frac{7}{11})(\frac{11}{11})(\frac{11}{24}) \]

\[ = \frac{7}{24}. \] (3.6)

**Example 2.**

Consider the same collection of 24 distinct low-lying configurations used in example 1. Look at the cluster \( (f_2 = B, f_3 = C, f_4 = D, f_5 = E) \) where \( B \) denotes the DNA marker \( S_{48} \), \( C \) denotes \( S_{46} \), \( D \) denotes \( S_4 \), \( E \) denotes \( S_{52} \).
For the above collection we find the frequency

\[ P_{2,3,4,5}(B, C, D, E) = \frac{12}{24}, \quad (3.7) \]

and we also find the frequencies

\[ P_{2,3,4}(B \mid C, D) = \frac{12}{15} \quad (3.8) \]
\[ P_{3,4,5}(C \mid D, E) = \frac{15}{16} \quad (3.9) \]
\[ P_{4,5}(D, E) = \frac{16}{24}, \quad (3.10) \]

again, the approximation (⋆) is satisfied by the observed frequencies.

However, since

\[ P_{2,3}(B \mid C) = \frac{13}{17} \quad (3.11) \]
\[ P_{3,4}(C \mid D) = \frac{15}{18} \quad (3.12) \]
\[ P_{4,5}(D \mid E) = \frac{16}{16} \quad (3.13) \]
\[ P_{2,3,4,5}(B, C, D, E) = \frac{12}{24}, \quad (3.14) \]

the frequencies do not satisfy the standard Markovian approximation.
\[ P_{2,3,4,5}(B, C, D, E) \neq P_{2,3}(B \mid C)P_{3,4}(C \mid D)P_{4,5}(D \mid E)P_5(E) \]

\[ 12/24 \neq (13/17)(15/18)(16/16)(16/24) \]

\[ 0.50 \neq 0.425 \quad (3.15) \]

A referee has kindly pointed out that in the context of Markov random fields (Spitzer [13]) one could perhaps find a rigorous basis for the observed Markov-like property. We thank the referee for this stimulating suggestion.

**IV. DISCUSSION**

The implementation of experimental techniques using RH data provides a bridge between chromosome mapping data generated from families and physical mapping data. RH experiments allow for the relative ordering of genetic markers that are too closely spaced to be detected by family linkage analysis. Additionally, it is not necessary to have polymorphic markers in order to generate useful information. Although not providing the level of resolution of physical mapping, RH mapping can complement and confirm results generated by pulse field gel electrophoresis.
Boehnke et al. [1] have presented a full discussion of the advantages and disadvantages of parametric and non-parametric ordering algorithms. As they point out, algorithms that search for minimum break configurations do not allow for estimates of distances between markers, nor do they provide relative likelihoods for one marker order over another. However, as the present work and the work of Boehnke et al. demonstrate, retention and inspection of sets of low-lying configurations yield important insight relating to the correlations of markers.

In our study we were interested in learning what properties might be present in a set of configurations with relatively few breaks. It became obvious that the set of low-lying configurations showed the clustering of particular markers. That was reassuring, since in complex optimization problems of the travelling salesman variety, one typically ends up with a set of low-lying configurations and never discovers the true absolute minimum.

One striking feature which we discovered here is that the set of low-lying configurations is described by a Markov-like probability distribution. That distribution contains all of the observed clustering of the markers. If one enlarges the set of low-lying configurations to include configurations with larger and larger numbers of breaks, the validity of the Markov-like approximation deteriorates. We are not in a position to judge whether the relation (*) is necessarily a deep or broad result, but we regard it as interesting.

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References

[1] M. Boehnke, K. Lange, D.R. Cox, Am. J.Hum. Genet. 49, 1174 (1991).
[2] D.R. Cox, M. Burmeister, E.R. Price, S. Kim and R.M. Myers, Science 250, 245 (1990).
[3] S.J. Goss and H. Harris, Nature 255, 680 (1975).
[4] S.J. Goss and H. Harris, J. Cell Sci. 25, 39 (1977).
[5] M. Burmeister, S. Kim, E.R. Price, T. de Lange, U. Tantravahi, R.M. Myers, and D.R. Cox, Genomics 9, 19 (1990).
[6] C.T. Falk, Genomics 9, 120 (1991).
[7] D.T. Bishop, G.P. Crockford, In: Genetic Analysis Workshop 7: Issues in Gene Mapping and Detection of Major Genes. J.W. MacCluer, A. Chakravarti, D.R. Cox, D.T. Bishop, S.J. Bale, M.H. Skolnick (eds.), Cytogenet Cell Genet 59, 93 (1992).
[8] D.E. Weeks, T. Lehner, J. Ott, In: Genetic Analysis Workshop 7: Issues in Gene Mapping and Detection of Major Genes. J.W. MacCluer, A. Chakravarti, D.R. Cox, D.T. Bishop, S.J. Bale, M.H. Skolnick (eds.), Cytogenet Cell Genet 59, 125 (1992).
[9] H. Falk, Physica, 104A, 459 (1980).
[10] T.M. Liggett, Interacting Particle Systems (Springer-Verlag, New York, 1985).
[11] S. Kirkpatrick, C.D. Gelatt and M.P. Vecchi, Science 220, 671 (1983).
[12] W.H. Press, B.P. Flannery, S.A. Teukolsky, and W.T. Vetterling, 
*Numerical Recipes: The Art of Scientific Computing* (Cambridge Univ. 
Press, London, 1986) pp. 326-334.

[13] F. Spitzer, *Random Fields and Interacting Particle Systems* 
(Mathematical Association of America, 1971); Am. Math. Monthly, 78, 
142 (1971).

Table 1.

Fourteen Chromosome-21 Markers Used in the Examples.

| APP | S1 | S4 | S8 | S11 | S12 | S16 | S18 | S46 | S47 | S47 | S52 | S111 | SOD |
|-----|----|----|----|-----|-----|-----|-----|-----|-----|-----|-----|------|-----|

Note: All numbered loci have a prefix of D21; APP = amyloid precursor; 
SOD = superoxide dismutase.
Table 2.

Minimum-Break Orders for Representative Simulations.

| Algorithm | Marker Order          |
|-----------|-----------------------|
| ↓ Breaks  |                       |
| 1 205     | APP S8 S1 S11 S16 S4 S12 S18 S48 S46 S52 S111 S47 SOD |
| 1 207     | APP S8 S1 S11 S16 S4 S12 S18 S46 S48 S52 S111 S47 SOD |
| 2 123     | S16 S48 S46 S4 S52 S11 S1 S18 S8 APP S111 S12 S47 SOD |
| 2 123     | S16 S48 S46 S4 S52 S11 S1 S18 S8 APP S111 S12 S47 SOD |
| 3 123     | S16 S48 S46 S4 S52 S11 S1 S18 S8 APP S111 S12 S47 SOD |
| 3 123     | S16 S48 S46 S4 S52 S11 S1 S18 S8 APP S111 S12 S47 SOD |

See note for Table 1.

Table 3.

24 Unique Marker Orders With Relatively Small Numbers of Breaks

| Breaks | Marker Order |
|--------|--------------|

23
| 123 | S16 | S48 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
|-----|-----|-----|-----|----|-----|-----|----|-----|----|-----|------|-----|-----|-----|
| 125 | S16 | S48 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S8  | S12 | S111 | S47 | SOD |
| 126 | S48 | S16 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 126 | S48 | S16 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
| 127 | S11 | S1 | S16 | S48 | S46 | S4 | S52 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
| 127 | S16 | S48 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 127 | S46 | S48 | S16 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 128 | S11 | S1 | S52 | S4 | S46 | S48 | S16 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 128 | S11 | S1 | S52 | S4 | S46 | S48 | S16 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
| 128 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S12 | S111 | APP | S8  | S18 | S11 | S11 |
| 128 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S111 | S12 | APP | S8  | S18 | S11 | S11 |
| 129 | S11 | S1 | S16 | S48 | S46 | S4 | S52 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 129 | S11 | S1 | S16 | S48 | S46 | S4 | S52 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
| 129 | S16 | S48 | S46 | S4 | S52 | S11 | S1 | S18 | S12 | S111 | S8 | APP | S47 | SOD |
| 129 | S16 | S48 | S46 | S4 | S52 | S11 | S1 | S18 | S12 | S111 | S8 | APP | S47 | SOD |
| 130 | S16 | S48 | S46 | S4 | S52 | APP | S8  | S12 | S111 | S47 | SOD | S18 | S1 | S11 |
| 130 | S16 | S48 | S46 | S4 | S52 | APP | S8  | S111 | S12 | S47 | SOD | S18 | S1 | S11 |
| 130 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | APP | S8  | S12 | S111 | S18 | S1 | S11 |
| 130 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S111 | S8 | APP | S18 | S1 | S11 |
| 130 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S111 | S8 | APP | S18 | S1 | S11 |
| 130 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S111 | App | S8  | S12 | S111 | S47 | SOD |
| 130 | S16 | S48 | S46 | S4 | S52 | SOD | S47 | S111 | S8 | APP | S12 | S111 | S47 | SOD |
| 130 | S48 | S16 | S46 | S4 | S52 | S11 | S1 | S18 | S8 | APP | S111 | S12 | S47 | SOD |
| 130 | S52 | S4 | S16 | S48 | S46 | S11 | S1 | S18 | S8 | APP | S12 | S111 | S47 | SOD |
| 130 | S52 | S4 | S46 | S48 | S16 | S11 | S1 | S18 | S8 | APP | S12 | S111 | S47 | SOD |

See note for Table 1.