Inferring Short-Range Linkage Information from Sequencing Chromatograms

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Abstract

Direct Sanger sequencing of viral genome populations yields multiple ambiguous sequence positions. It is not straightforward to derive linkage information from sequencing chromatograms, which in turn hampers the correct interpretation of the sequence data. We present a method for determining the variants existing in a viral quasispecies in the case of two nearby ambiguous sequence positions by exploiting the effect of sequence context-dependent incorporation of dideoxynucleotides. The computational model was trained on data from sequencing chromatograms of clonal variants and was evaluated on two test sets of in vitro mixtures. The approach achieved high accuracies in identifying the mixture components of 97.4% on a test set in which the positions to be analyzed are only one base apart from each other, and of 84.5% on a test set in which the ambiguous positions are separated by three bases. In silico experiments suggest two major limitations of our approach in terms of accuracy. First, due to a basic limitation of Sanger sequencing, it is not possible to reliably detect minor variants with a relative frequency of no more than 10%. Second, the model cannot distinguish between mixtures of two or four clonal variants, if one of two sets of linear constraints is fulfilled. Furthermore, the approach requires repetitive sequencing of all variants that might be present in the mixture to be analyzed. Nevertheless, the effectiveness of our method on the two in vitro test sets shows that short-range linkage information of two ambiguous sequence positions can be inferred from Sanger sequencing chromatograms without any further assumptions on the mixture composition. Additionally, our model provides new insights into the established and widely used Sanger sequencing technology. The source code of our method is made available at http://bioinf.mpi-inf.mpg.de/publications/beggel/linkageinformation.zip.

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Introduction

Direct Sanger sequencing is cheap and widely used but suffers from low sensitivity regarding the detection of minor variants and from the loss of linkage information. The problem of reconstructing the linkage of several ambiguous sequence positions has been addressed only in the setting of diploid genomes so far. Linkage information is of clinical relevance as treatment decisions, for instance for human immunodeficiency virus and hepatitis B patients, are often based on Sanger sequence data of the viral genome [1–3]. A particular problem arises when two ambiguous sequence positions are present within the same codon (Figure 1). Then the amino acids expressed cannot be determined precisely due to the lack of linkage information.

Significant attempts have been made to infer linkage information from Sanger sequencing data [4–9]. These methods are tailored to genomes that contain a mixture of two different sequences one of which harbors an insertion or deletion. In this case, the chromatogram downstream of the insertion or deletion shows a high number of double peaks as the two genetic variants are superimposed with a phase shift. These methods rely on prior knowledge (reference sequence or set of possible single nucleotide polymorphisms (SNP)) or the availability of both the forward and the reverse chromatogram. Otherwise, such methods can only be applied in situations where the two mixture sequences are sufficiently similar and the analyzed fragment is significantly longer than the insertion or deletion [4]. The method we present here does not rely on diploid genomes that contain a heterozygous insertion or deletion. It is tailored to viral genomes that can have a more complex quasispecies without insertions or deletions. Additionally, we explore a source of information that is encoded in the peak heights of the sequencing chromatograms and has not been considered so far.

Dye-labeled terminator sequencing chromatograms exhibit heterogeneous peak heights that result from different affinities of the polymerase for dideoxynucleotides (chain terminator nucleotides) rather than natural nucleotides during polymerase chain reaction amplification [9–11]. The rate of incorporation has been found to also depend on the up- and downstream subsequence [12,13]. This effect is referred to as sequence context-dependent incorporation of dideoxynucleotides. Methods for estimating the relative frequency of two DNA bases present at a sequence position account for this effect by comparing the observed peak heights to the peak heights of reference chromatograms [13,14].
Interaction effects between peaks of nearby ambiguous sequence positions are considered to be confounding the interpretation of chromatograms and, therefore, are disregarded or averaged out by these methods. In contrast, we have found that such interactions harbor valuable information for inferring haplotype frequencies. We have developed a probabilistic model that exploits this interaction for the purpose of reconstructing the linkage and thereby also determines the amino acids expressed at two nearby positions.

The key assumption of our approach is that the collection of peak heights of a mixture of clonal variants (which are called haplotypes, in the following) is the proportion-weighted mixture of the peak heights of the underlying haplotypes. The underlying physico-chemical reasoning is as follows. Context-dependent incorporation of dideoxynucleotides resulting in context-dependent peak heights occurs at the level of each polymerase molecule that incorporates dideoxynucleotides with an affinity depending on the DNA sequence it is processing. The observed sequencing chromatogram is the sum of all individual molecular fluorescence impulses. Thus knowing the peak heights of all possible haplotypes present in a mixture, given that these show distinct patterns, renders the determination of the haplotype composition feasible. As context-dependent incorporation of dideoxynucleotides is a local effect the methodology we present here will only infer linkage information over a short genomic region, maybe up to 5 bases.

Data preparation for the preliminary analysis

The chromatograms of three dilution series published in [14] were obtained from the authors. Each dilution series was created by mixing different pairs of DNA fragments that differ at a single genomic position (single-nucleotide polymorphism). Experimental mixture proportions were 10:0, 9:1, 8:2, 7:3, 6:4, 5:5, 4:6, 3:7, 2:8, 1:9 and 0:10. Thus, eleven samples per dilution series were available. Each sample was amplified and sequenced three times.

In vitro experimental setup

The wild-type HBV plasmid vector pCH9-3091 was modified by site-directed mutagenesis. This resulted in a set of six clonal HBV variants (Table 1). The clonal variants v1,v2,v3,v4 represent all possible haplotypes for the two ambiguous positions 610 and 612 in the Reverse Transcriptase shown in Figure 1. Likewise, the haplotypes v1,v2,v3,v6 represent all possible haplotypes for two SNPs at positions 610 and 614. Test set 1 (TS1) consists of 29 in vitro mixtures of the four haplotypes v1,v2,v3,v4 and test set 2 (TS2) consists of 42 in vitro mixtures of the four haplotypes v1,v2,v3,v6. Mixtures were prepared by equimolar mixing. The in vitro mixtures were submitted to independent sequencing reactions, each with the same set of oligonucleotides. The compositions of the test sets are available in Table S1.

Materials and Methods

In a preliminary study we analyzed a set of sequencing chromatograms from three dilution series published in [14]. This motivated the mixture assumption and provided evidence that sequencing chromatograms contain linkage information and that reconstructing the haplotype composition might be feasible. In order to evaluate our hypothesis we designed and created two in vitro test sets, each consisting of four clonal hepatitis B virus (HBV) variants that differ at two nucleotide positions along with two sets of in vitro mixtures of the four respective haplotypes.
Table 1. Experimental setup.

| Haplotype | Position 610 | Position 612 | Position 614 | Comment |
|-----------|--------------|--------------|--------------|---------|
| v1        | Adenine      | Guanine      | Adenine      | Wild-type |
| v2        | Guanine      | Guanine      | Adenine      | rtM204V  |
| v3        | Adenine      | Thymine      | Adenine      | rtM204I  |
| v4        | Guanine      | Thymine      | Adenine      | rtM204V  |
| v5        | Adenine      | Guanine      | Thymine      |          |
| v6        | Guanine      | Guanine      | Thymine      |          |

Six clonal hepatitis B virus genomes (haplotypes) were created by varying three nearby nucleotide positions. Haplotypes v1, v2, v3, v4, v5, v6 represent all possible combinations of the two variants adenine and guanine present at position 610 and the two variants guanine and thymine present at position 612. These nucleotide variants are clinically relevant as they represent two primary resistance mutations rtM204I and rtM204V. Haplotypes v1, v2, v3, v4 represent all possible combinations of the two variants adenine and guanine present at position 610 and the two variants adenine and thymine present at position 614. Positions are numbered with respect to the Reverse Transcriptase of the hepatitis B virus genome.

Conditional distribution assumption

The computational approach requires repetitive sequencing of all possible clonal variants in the mixture as training data. Each haplotype has its own characteristic sequencing chromatogram profile because of the effect of context-dependent incorporation of dideoxynucleotides. The chromatogram of a mixture of haplotypes is then the proportion weighted combination of these four haplotype-specific profiles (with some variance). As a preprocessing step all sequencing chromatograms were normalized using a sequence region (called the normalization region) upstream from the region under analysis. Each DNA base (A, T, C, G) requires its own normalization parameter. These four parameters were set to reflect sequencing-dependent variations in peak heights, which were estimated using sequencing replicates of clonal variants.

Data likelihood

In order to apply equation (3) (to compute the data likelihood \( P(D|M,z) \)) the normalization constants \( \gamma_{B|h}\) need to be estimated. Therefore, the data D is partitioned into the distinct union of \( D= D_{\text{eval}} \cup D_{\text{fit}} \). \( D_{\text{eval}} \) contains all ambiguous and \( D_{\text{fit}} \) the remaining peak heights. The normalization constants \( \gamma_{B|h}\) are estimated on \( D_{\text{fit}} \) using the maximum likelihood principle given the conditional distribution assumption (3) and the mixture profile \( m_{i} \). Fitting each normalization constant is an ordinary least-square regression problem for each mixture profile \( m_{i} \). The likelihood of the observation \( P(D|M,z) \) is then computed using equation (4). Thus, we evaluate \( P(D|M,z) \) by computing \( P(D_{\text{eval}}|M,z,D_{\text{fit}}) \). After the normalization constants \( \gamma_{B|h}\) have been estimated using \( D_{\text{fit}} \) the peak heights of \( D_{\text{eval}} \) can be assumed to be independent, given the conditional distribution assumption (3). This allows the likelihood function to be factorized.

\[
P(D|M,z) = P(D_{\text{eval}}|M,z,D_{\text{fit}}) = \prod_{h \in D_{\text{eval}}} P(\gamma_{B|h}|M,z) (4)
\]

Model selection

The primary goal is to determine the haplotypes \( M \subseteq \{1,\ldots,4\} \) represented by a given chromatogram. Maximum likelihood estimates derived from the conditional distribution assumption overfit and likely produce point estimates indicating that all four haplotypes are present. In order to guard against overfitting regularization is employed by computing the marginal model likelihoods according to equation (5) with uniform priors \( P(z|M) \) [17].

\[
P(D|M) = P(D_{\text{eval}}|M,D_{\text{fit}}) = \int P(D_{\text{eval}}|z,D_{\text{fit}}) P(z|M) dz (5)
\]

For the setting with two ambiguous sequence positions seven alternative models need to be considered, which are \( M_{1+4}, M_{2+3}, M_{1+2+3}, M_{1+2+4}, M_{1+3+4}, M_{2+3+4} \) and \( M_{1+2+3+4} \). Thus, inferring the haplotype composition is a 7-fold classification problem. The notation \( M_{1+4} \) is used to indicate a mixture of haplotypes 1 and 4, \( M_{1+2+3} \) is used to indicate a mixture of haplotypes 1, 2 and 3, etc.. Other subsets of \( \{1,\ldots,4\} \), e.g. \( \{1,2\} \), do not exhibit two ambiguous positions in the chromatogram and therefore can be excluded.

Additionally, the modes \( \hat{\beta}_{1}, \ldots, \hat{\beta}_{4} \) of the posterior marginals of the model parameters \( z_{1}, \ldots, z_{4} \) of the full model \( M_{1+2+3+4} \) are computed. For each \( j \in \{1,2,3,4\} \) let \( \hat{z}_{j} \) denote all \( z_{i} \) with \( i \neq j \).
fulfilled if the model under consideration is not the full model parameter marginals. The latter condition can and need only to be together indicate a strong contradiction of the model with the preliminary analysis

Performance Evaluation

and a haplotype haplotypes (Table 1). Further, heights upstream of the ambiguous sequence position were almost ambiguous sequence position and found that the normalized peak heights of the underlying haplotypes. The mixture assumption is common-sense for the ambiguous sequence position in a dilution series. Methods for quantifying base frequencies based on chromatograms rely on this principle [13, 14]. Additionally, we found that this holds for all peak heights. It becomes evident near ambiguous sequence positions where the peak heights of the underlying haplotypes differ due to the effect of sequence context-dependent incorporation of dideoxynucleotides. Finally, Figure 2 shows that there is indeed linkage information in sequencing chromatograms as the peak height at e.g. position 2 depends on the relative frequencies of the bases present at the ambiguous sequence position.

The mixture assumption was employed to estimate the mixture fractions of three dilution series. The maximum likelihood estimates for the mixture fractions derived from the chromatograms were close to the nominal mixture fractions for dilutions series 1 and 2 with average absolute errors of 0.01 and 0.03, respectively (Figures 3A and 3B). Dilution series 3 showed a nonlinear relationship between nominal and estimated proportions, resulting in an average absolute error of 0.14 (Figure 3C). This nonlinear relationship is likely caused by varying amplification efficiencies of the different clonal variants. Almost identical results for these data were reported by [14]. Note that the variance in the estimates originates from repetitive amplification and sequencing of the same in vitro mixture. The experiment described above was repeated while blanking out the ambiguous positions. Thus, we employed only sequence positions $-2, -1, 1$ and 2 (Figure 2) to estimate the mixture fractions. This resulted in fraction estimates with higher average errors of 0.05, 0.08, 0.26 and higher variances (Figures 3D–3E). The mixture assumption facilitates the estimation of the fractions of a mixture without using the ambiguous positions. Previous methods that quantify mixture fractions based on sequencing chromatograms only consider ambiguous positions and neglect the information provided by surrounding peaks [13, 14].

We now focus on the problem of haplotype reconstruction based on sequencing chromatograms in the special case of two ambiguous sequence positions. For this problem four possible haplotypes $1, 1, 4$ with fractions $z_1, z_2, z_3$ need to be considered. Table 1 rows 1 to 4 lists the four possible haplotypes for the combination of two important HBV drug resistance mutations within the Reverse Transcriptase domain. The haplotype reconstruction problem has three degrees of freedom $z_1, z_2, z_3$ which determine $z_4$ due to $\sum z_i = 1$. An estimate of the fraction of the first ambiguous position $p_1$ gives rise to $z_1 + z_2 = p_1$ as haplotype 1 and 3 are wild-type at the first ambiguous position. Similarly, an estimate $p_2$ of the fraction of the second ambiguous position provides $z_1 + z_2 = p_2$. These frequency estimates can be derived from the chromatograms thus reducing the complexity to one degree of freedom. This implies that the peak heights of the ambiguous positions impose strong constraints that limit the space of possible solutions to a straight line in the three-dimensional space of all possible solutions. Within this set of possible solutions we face the problem of model selection. Further evidence from the surrounding peaks or slightly different configurations of the four ambiguous peaks are required to localize the correct solution.

$\beta_i = \arg \max_{\beta_i} P(x|D, M_{1+t+3+4})$

$= \arg \max_{\beta_j} \int_{[0,1]^3} P(x|D, M_{1+t+3+4}) \, dx \, d\beta_j$

$\sim \arg \max_{\beta_j} \int_{[0,1]^3} P(x|M_{1+t+3+4}) P(D|M_{1+t+3+4}) \, dx \, d\beta_j$

A model $M \subseteq \{1,\ldots,4\}$ is considered to be inconsistent with $\beta_1, \ldots, \beta_4$ given a certain cutoff value $t$ if there exists $j \in M$ with $\beta_j \leq t$ and (in case $M \neq M_{1+t+3+4}$) if there is $k \notin M$ with $\beta_k > t$. In words, there is a haplotype $j$ indicated by the model with low $\beta_j$ and a haplotype $k$ not present in the model with high $\beta_k$, which together indicate a strong contradiction of the model with the parameter marginals. The latter condition can and need only to be fulfilled if the model under consideration is not the full model $(M_{1+t+3+4})$. Models that are inconsistent with the modes of the parameter marginals are excluded from the model selection procedure. The cutoff $t$ used to evaluate the prediction performance on TS1 was chosen to optimize accuracy on TS2 and vice versa.

The respective integrals (5) and (6) were approximated using a grid of $x$ values of precision 0.025. This resulted in 12341 parameter configurations, for which the log-likelihoods were computed and summed up to compute the marginal model likelihoods and the posterior marginal parameter distributions.

Performance Evaluation

Haplotype reconstruction for two ambiguous sequence positions is a 7-fold classification problem. Prediction performance at the model level is measured in terms of accuracy. Prediction performance is also evaluated at the clonal level. For this purpose, each 7-fold classification problem is interpreted as four 2-fold performance is also evaluated at the clonal level. For this purpose, each $7 \times 7$ classification problem is interpreted as four $2 \times 2$ fractions based on sequencing chromatograms in the special case of two ambiguous positions. Previous methods that quantify mixture fractions based on sequencing chromatograms only consider ambiguous positions and neglect the information provided by surrounding peaks. The respective integrals (5) and (6) were approximated using a grid of $x$ values of precision 0.025. This resulted in 12341 parameter configurations, for which the log-likelihoods were computed and summed up to compute the marginal model likelihoods and the posterior marginal parameter distributions.

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Results

We developed a computational model to describe the expected sequencing chromatogram of a mixture of haplotypes. Such a model facilitates the inference of the haplotype composition i.e. to infer the mixture components and fractions. To validate the model two test sets were created. Each test set consists of four clonal HBV variants, which differ at two nearby nucleotide positions, and corresponding test samples of in vitro mixtures of the respective haplotypes (Table 1). Further, in silico test samples were created to study limitations of our approach.

Preliminary analysis

We analyzed chromatograms of a dilution series with nominal mixture proportions 10:0, 9:1, 8:2, 7:3, 6:4, 5:5, 4:6, 3:7, 2:8, 1:9 and 0:10. We were interested in changes of peak heights near the ambiguous sequence position and found that the normalized peak heights upstream of the ambiguous sequence position were almost identical for all nominal mixture proportions. At the ambiguous sequence position and at the two to four bases downstream we observed a smooth and almost linear transition of the peak heights between the samples with nominal mixture proportions 10:0 and 0:10 (Figure 2). This gave rise to the mixture assumption: the peak heights of a mixture of haplotypes are the proportion-weighted mixtures of the peak heights of the underlying haplotypes. The mixture assumption is common-sense for the ambiguous sequence position in a dilution series. Methods for quantifying base frequencies based on chromatograms rely on this principle [13, 14]. Additionally, we found that this holds for all peak heights. It becomes evident near ambiguous sequence positions where the peak heights of the underlying haplotypes differ due to the effect of sequence context-dependent incorporation of dideoxynucleotides. Finally, Figure 2 shows that there is indeed linkage information in sequencing chromatograms as the peak height at e.g. position 2 depends on the relative frequencies of the bases present at the ambiguous sequence position.

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In silico experiment

In order to study elementary properties of our approach we created a set of in silico test samples. 1771 artificial chromatograms were created for the set of haplotypes $v_1, \ldots, v_4$ (see Table 1) based on equation (2) using a grid of $z$ values with grid size 0.05. Test data were generated without adding a noise component. The noise-free setting provides an upper bound on the model performance to be expected on real-world data and simplifies the analysis of model characteristics and limitations. Mixture chromatograms were truncated to nucleotide positions 600 to 620. Positions 610 and 612 were employed to compute the data likelihoods. The value of the standard deviation $\sigma$ was estimated using the sequencing repetitions of the haplotypes. $\sigma$ was set to 20.94.

Only 867 (49%) of the 1771 in silico test samples were predicted correctly. Figure 4 visualizes all incorrectly predicted test cases separately for each falsely predicted label. The four surfaces of the simplex correspond to the four 3-mixture models $M_{123}$, $M_{124}$, $M_{134}$ and $M_{234}$. Test samples that lie in the interior of the simplex but close to one of its faces display mixtures of all four haplotypes with one haplotype having low frequency (Figures 4A–4D). These test samples were regularized towards the surface by using the marginal likelihoods for model selection. This reveals a major restriction of the model in terms of sensitivity.

Figure 2. Peak heights of dilution series. The figure shows the median normalized peak heights of the chromatograms of a dilution series sorted by nominal mixture proportion. The normalized peak heights before the ambiguous sequence position are almost identical for all nominal mixture proportions. At the ambiguous sequence position and at up to five bases downstream of the ambiguous sequence position a smooth and apparently linear transition between the peak heights of the samples with nominal mixture proportions 10:0 and 0:10 can be observed.

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Figure 3. Fraction estimates for dilution series. All subplots show nominal mixture fractions versus estimated mixture fractions for three dilution series. A–C use all peak heights and provide proportions estimates with low error and low variance. D–E ignore the peak heights at the ambiguous positions and try to estimate the mixture proportions based on the unambiguous positions only. The resulting fraction estimates show higher error and higher variance.

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The test set can be divided into two groups: those samples that contain at least one haplotype with a fraction of no more than 10%, and those for which each present haplotype occurs in a fraction higher than 10%. The samples of the first group were predicted correctly in 303 (32.0%) of 1200 cases. The samples of the second group (all present haplotypes exceed a fraction of 10%) were predicted correctly in 404 (85%) of 571 cases.

Figure 4E indicates a second systematic failure: 34 \( M_{1+2+3+4} \), 5 \( M_{1+2+3} \) and 7 \( M_{1+3+4} \) samples were incorrectly predicted as mixtures of haplotypes 2 and 4. All of these samples satisfied the set of linear constraints \( x_2 - x_3 \leq 0.05 \), \( x_2 \leq 0.35 \) and \( x_3 \leq 0.35 \). Similarly, 32 \( M_{1+2+3+4} \), 5 \( M_{1+2+4} \) and 4 \( M_{1+3+4} \) samples were falsely predicted as mixtures of haplotypes 2 and 3 (Figure 4F). These samples satisfied the constraints \( 2x_1 + x_2 + x_3 - 1 \leq 0.05 \), \( x_1 \leq 0.30 \) and \( x_2 \leq 0.30 \). In order to improve prediction performance we analyzed the marginal posterior distributions of the fraction parameters \( x_1, \ldots, x_4 \). E.g. for all of the seven \( M_{2+3+4} \) test samples that were misclassified as \( M_{1+4} \) we observed that the modes of posterior marginal distributions of \( x_1 \) were zero while the modes of \( x_2 \) had a mean of 0.22±0.07. Thus, the modes of the posterior marginals were clearly inconsistent with the model predictions based on the marginal model likelihoods. By checking the consistency of the model predictions with the modes of the posterior marginals using a cutoff-value of 10% (as described in the Methods section) all misclassified 3-mixture test samples could be corrected. Nevertheless, the consistency rule does not improve prediction performance on the 4-mixture test samples. The inability of the model to distinguish between 2-mixture and 4-mixture test samples (if one of two sets of linear constraints is fulfilled) is a major limitation of the model in terms of accuracy.

**In vitro validation**

Our hypothesis was that the haplotype composition of arbitrary mixtures can be reconstructed using information from the ambiguous and the surrounding peak heights. In order to test this hypothesis, we created two in vitro training sets, each composed of six sequencing repetitions of the underlying haplotypes (Table 1) to compute the individual haplotype profiles and a respective test set of in vitro mixtures. Test set 1 (TS1) based on haplotypes \( v_1, v_2, v_3, v_4, v_5 \) contained 29 mixtures and test set 2 (TS2) based on haplotypes \( v_1, v_2, v_3, v_6 \) contained 42 mixtures. When model selection was solely based on the marginal model likelihoods prediction accuracy at the model level was 96.6% on TS1 and 47.6% on TS2. By application of the consistency rule as described in the Methods section prediction accuracy improved to 71.4% on TS2 and remained unchanged on TS1. The respective cutoff used to evaluate the prediction performance on TS1 was chosen to optimize accuracy on TS2 and vice versa. A cutoff of 10% was used for both data sets. Note that model selection in our setting is a 7-fold classification problem and the accuracy expected by chance amounts to only 14.3%. Prediction performance was also evaluated at the clonal level. In this evaluation scheme we treated the 7-fold classification problem as four 2-fold classification problems that amounted to the prediction of the presence or the absence of each of the four possible haplotypes. Using this evaluation we could study the accuracy of the model predictions in more detail than when assessing accuracy in terms of 0–1 error at the model level. For instance, the incorrect classification of a \( M_{1+2+3+4} \) test sample as \( M_{1+2+3} \) would result in an accuracy of 75% as three of four haplotypes were predicted correctly. Clonal level prediction accuracy was 97.4% for TS1 and 84.5% for TS2. Figure 5 summarizes the prediction performances on TS1 and TS2 both at the model level and at the clonal level. Additionally,

**Figure 4. In silico prediction results.** 1771 in silico test chromatograms were created by computing the mixture profiles on a grid of \( s \) values with precision 0.05. Test chromatograms were classified by the mixture model with \( s = 0.94 \). The subplots show all falsely classified samples separately for each falsely predicted label. Six major cases of misclassification can be observed. Subplots A–D show test samples that consist of four haplotypes with at least one haplotype having low frequency. Subplots E and F show test samples that were predicted as mixtures of haplotypes 1 and 4 or of haplotypes 2 and 3, respectively. The data points of subplot E satisfy the linear constraints \( x_2 - x_3 \leq 0.05 \), \( x_1 \leq 0.30 \) and \( x_4 \leq 0.30 \). The data points of subplot F satisfy \( 2x_1 + x_2 + x_3 - 1 \leq 0.05 \), \( x_1 \leq 0.30 \) and \( x_2 \leq 0.30 \). DOI:10.1371/journal.pone.0081687.g004
we present the results as a function of the prediction confidence expressed by the uncertainty cutoff. E.g. at an uncertainty cutoff of 4.0 only 2.6% (5.4%) of predictions at the clonal level were incorrect, 14.6% (30.4%) were unassigned and 82.8% (64.2%) of predictions were correct on TS1 (respectively TS2).

Discussion

This article describes the first attempt, to our knowledge, to use peak heights from sequencing chromatograms to infer linkage information about nearby ambiguous positions. We derived a generative model to describe the expected peak heights of a mixture of haplotypes by combining the chromatograms of the underlying haplotypes. The model was motivated by visual inspection of the peak heights of a dilution series. Our model is a generalization of previous methods for estimating the fractions of one ambiguous sequence position [13,14]. The main difference is that we employ a generative model whose parameters are estimated by probabilistic inference rather than by comparing the observed peak heights to the template peak heights. Previous methods neglected nearby peak heights as they are usually influenced by the ambiguous peaks to be analyzed. In contrast, we exploit this interaction effect to infer linkage information.

The high prediction accuracy of our method on two in vitro test sets of 29 and 42 mixture samples shows that sequencing chromatograms do contain linkage information. The prediction accuracy on TS1 was 97.4% at the clonal level and 96.6% at the model level while the accuracy on TS2 was 84.5% at the clonal level and 96.6% at the model level. We found that the marginal likelihood of the best model divided by the marginal likelihoods of all other models falls below the uncertainty cutoff displayed on the x-axis.

The evaluation of the model on a set of 1771 in silico test samples revealed several limitations of our approach. First, the model cannot reliably detect minor populations with frequency of no more than 10%. In such cases, the predictions are regularized towards simpler models, which do not contain the low-frequency haplotype. Second, we found that if the mixture fractions \( z_j \) satisfy one of two sets of linear constraints then the mixture is always predicted to be either a mixture of the haplotypes 1 and 4 or to be a mixture of haplotypes 2 and 3. The true mixture components in the former case are \( M_{1+4} \), \( M_{1+2+3} \), \( M_{2+3+4} \) and \( M_{1+2+3+4} \) and in the latter case are \( M_{2+3} \), \( M_{1+2+4} \), \( M_{1+3+4} \) and \( M_{1+2+3+4} \).

Figure 5. Prediction accuracy on in vitro test sets. The figure shows the prediction accuracy on test sets TS1 (subplots A and B) and TS2 (subplots C and D). Prediction accuracy was evaluated both at the clone and at the model level. Each test sample is either predicted correctly, predicted incorrectly or unassigned. The latter happens when the marginal likelihood of the best model divided by the marginal likelihoods of all other models falls below the uncertainty cutoff displayed on the x-axis.

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We evaluated our hypothesis to perform inference on the haplotype compositions for only two sets of sequence positions within the Reverse Transcriptase domain of the HBV genome. Nevertheless, peak heights usually display very distinctive patterns and thus we assume the approach will also work in other settings. Two different amplification and sequencing protocols in concert with two different sequencing machines were employed to generate the two test sets. This indicates the robustness of the method with respect to sequencing protocols. The reasoning that the problem is essentially one-dimensional after the mixture fractions are estimated implies that the model very likely cannot successfully be extended to infer the haplotype composition in the presence of more than two nearby ambiguities. Additionally, the locality of the effect of sequence context-dependent incorporation of dideoxynucleotides limits the capacity of the approach to infer phase information of ambiguous positions that are more than a few bases (up to five bases, at most) apart from each other. On test set TS2, in which the two ambiguous positions were three positions apart from each other, prediction accuracy was impaired in comparison to TS1, in which the two ambiguous positions were only separated by a single base.

The advent of next-generation sequencing (NGS) technologies has superseded traditional Sanger sequencing in many applications [18]. NGS technologies offer increased sequencing depth and speed through a high degree of parallelization and miniaturization and allow the detection of minor variants with relative frequencies of as low as 10^{-4} [19]. NGS data also naturally provide linkage information over the whole read length, which may further be extended by the use of paired-end reads [20]. Compared to our approach, NGS technologies are far more sensitive and accurate in determining the haplotype composition of a mixture and can deliver long-range linkage. Nevertheless, traditional Sanger sequencing is cheaper than NGS and NGS will not be accessible in many laboratories for some time to come. Before this background, certain applications, in which the limitations of our approach in terms of sensitivity, accuracy and limited range are acceptable, may not require NGS.

Conclusions

We have developed and validated an approach to compute the peak heights of a mixture of haplotypes based on the chromatograms of the underlying haplotypes. The model can be used to infer the haplotypes present in a mixture and therefore the short-range linkage for two ambiguous sequence positions. The effect of sequence context-dependent incorporation of dideoxynucleotides — an effect that was previously regarded as detrimental — was employed. The effectiveness of our method shows that short-range linkage information can be inferred from sequencing chromatograms with no further assumptions on the mixture composition. The model also allows the estimation of the fractions of nearby ambiguities and therefore overcomes the limitations of [1]. As a major limitation to its widespread applicability the model requires the sequencing chromatograms of all possible haplotypes in the mixture. The source code of our method can be downloaded at http://bioinf.mpi-inf.mpg.de/publications/beggel/linkageinformation.zip.

Supporting Information

Table S1 Composition of the in vitro test set 1 (TS1) and test set 2 (TS2).

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Author Contributions

Conceived and designed the experiments: BB MNF JV RK. Performed the experiments: MNF. Analyzed the data: BB. Contributed reagents/materials/analysis tools: MNF. Wrote the paper: BB TL.

References

1. Lengauer T, Sing T (2006) Bioinformatics-assisted anti-HIV therapy. Nat Rev Microbiol 4: 790–7.
2. Lengauer T, Sander O, Sierra S, Thielen A, Kaiser R (2007) Bioinformatics prediction of HIV coreceptor usage. Nat Biotechnol 25: 1807–10.
3. Zoulim F, Durantel D, Deny P (2009) Management and prevention of drug resistance in chronic hepatitis B. Liver Int 29 Suppl 1: 108–15.
4. Dmitriev DA, Raktov RA (2008) Decoding of superimposed traces produced by direct sequencing of heterozygous indels. PLoS Comput Biol 4: e1000115.
5. Plot JF, Tillier A, Samadi S, Tillier S (2006) Phase determination from direct sequencing of lengthvariable DNA regions. Molecular Ecology Notes 6: 627–630.
6. Plot JF (2007) champuru 1.0: a computer software for unraveling mixtures of two DNA sequences of unequal lengths. Molecular Ecology Notes 7: 974–977.
7. Sousa Santos C, Rhabo JL, Colares-Pereira MJ, Almada VC (2005) Heterozygous indels as useful tools in the reconstruction of DNA sequences and in the assessment of ploidy level and genomic constitution of hybrid organisms. DNA Seq 16: 462–7.
8. Seroussi Y, Seroussi E (2007) TraceHaplotyper: using direct sequencing to determine the phase of an indel followed by biallelic SNPs. Biotechniques 43: 455–458.
9. Li Y, Mitawko V, Waksman G (1999) Structure-based design of Taq DNA polymerases with improved properties of dideoxynucleotides. Proc Natl Acad Sci USA 96: 9491–6.
10. Tabor S, Richardson CC (1995) A single residue in DNA polymerases of the Escherichia coli DNA polymerase I family is critical for distinguishing between deoxy- and dideoxyribonucleotides. Proc Natl Acad Sci USA 92: 6339–43.
11. Lee LG, Connell CR, Woo SL, Cheng RD, McArule BF, et al. (1992) DNA sequencing with dye-labeled terminators and T7 DNA polymerase: effect of dyes and dNTPs on incorporation of dye-terminators and probability analysis of termination fragments. Nucleic Acids Res 20: 2471–83.
12. Parker LT, Zakeri H, Deng Q, Spurgeon S, Kwok PY, et al. (1996) AmpliTaq DNA polymerase, FS dye-terminator sequencing: analysis of peak height patterns. Biotechniques 21: 694–9.
13. Kwok PY, Carlson C, Yager TD, Ankener W, Nickerson DA (1994) Comparative analysis of human DNA variations by fluorescence-based sequencing of PCR products. Genomics 23: 138–44.
14. Curr IM, Robinson JJ, Diminou R, Markham AF, Morgan AW, et al. (2009) Inferring relative proportions of DNA variants from sequencing electropherograms. Bioinformatics 25: 3244–50.
15. Schildgen O, Schewe CK, Vogel M, Däumer M, Kaiser R, et al. (2004) Successful therapy of hepatitis B with tenofovir in HIV-infected patients failing previous adefovir and lamivudine treatment. AIDS 18: 2325–7.
16. Zhang Q, Wu G, Richards E, Jia S, Zeng C (2007) Universal primers for HBV genome DNA amplification across subtypes: a case study for designing more effective viral primers. Virol J 4: 92.
17. Barber D (2012) Bayesian Reasoning and Machine Learning. Cambridge: Cambridge University Press.
18. Metzker ML (2010) Sequencing technologies - the next generation. Nat Rev Genet 11: 31–46.
19. Gerstung M, Brüdel C, Rechsteiner M, Wild P, Schraml P, et al. (2012) Reliable detection of subclonal single-nucleotide variants in tumour cell populations. Nat Commun 3: 811.
20. Chaisson MJ, Brinza D, Pevzner PA (2009) De novo fragment assembly with short mate-paired reads: Does the read length matter? Genome Res 19: 336–46.