

1 Collapsed Haplotype Pattern Method for Linkage Analysis of Next-Generation Sequence Data

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Abstract

Recent advances in next generation sequencing (NGS) make it possible to directly sequence genomes and exomes of individuals with Mendelian diseases and screen sequence data for causal variants. With the reduction in cost of NGS, DNA samples from entire families can be sequenced and linkage analysis can be performed directly using NGS data. Inspired by “burden” tests which are used for complex trait rare variant association studies, we developed the collapsed haplotype pattern (CHP) method for linkage analysis. Using data from several deafness genes we demonstrate that the CHP method is substantially more powerful than analyzing individual variants. Unlike applying NGS data filtering approaches, the CHP method provides statistical evidence of a gene’s involvement in disease etiology and is also less likely to exclude causal variants in presence of phenocopies and/or reduced penetrance. The CHP method was implemented in the SEQLinkage software package which can perform linkage analysis on NGS data or can generate data compatible with many linkage analysis programs, reviving them for use in NGS era.

Keywords

Parametric linkage analysis, LOD scores, Mendelian diseases and traits, Next-generation sequence data, Statistical method.

32 **Introduction**

33 The advent and advance of NGS in recent years has led to the identification of a large number of
34 Mendelian disease genes. The typical approach to identifying Mendelian disease causal variants
35 using either whole genome sequence (WGS) or exome sequence (WES) data is to filter variants in
36 an affected individual or shared by affected family members, excluding those which are found at
37 higher frequencies, e.g. $>0.5\%$ in variant databases. Sometimes unaffected family member(s) are
38 also used in the filtering process. While filtering is straightforward and has been successful¹, such
39 efforts rely on limited family information, e.g. mode of inheritance, sharing between a subset of
40 family members and information from external resources on variant functional characterizations
41 and frequencies. On the other hand, linkage analysis, which incorporates information on mode of
42 inheritance, penetrance, allele frequencies and genetic map information, remains a powerful tool
43 to localize Mendelian disease loci. As a result, combined SNP array based linkage analysis and
44 sequence based filtering method is becoming popular². There is also a great interest to directly
45 perform linkage analysis on rare variants obtained from NGS data. Although it has been shown
46 that analyzing rare single nucleotide variants (SNVs), usually designated as having a minor allele
47 frequency (MAF) $<0.5\%$ or 1% , from NGS data provides acceptable linkage results, due to low
48 heterozygosity of SNVs and allelic heterogeneity this approach can be less powerful than analysis
49 of SNPs from genotyping arrays³.

50 Here we describe the collapsed haplotype pattern (CHP) method which is motivated by rare variant
51 association methods which analyze multiple rare variants within a region, which is often a gene.
52 The CHP method was designed to analyze rare variants by constructing markers that have a higher

heterozygosity and are more informative for linkage analysis than individual rare SNVs. Unlike when SNPs are analyzed, the CHP method does not require linkage disequilibrium (LD) pruning to avoid spurious associations⁴. The CHP method is particularly powerful in the presence of intra- (e.g. compound heterozygotes) and inter-family allelic heterogeneity, a phenomenon commonly observed for Mendelian diseases. When causal variants are missing from samples, the CHP method can still detect linkage due to transmission information retained by other variants. We have developed the SEQLinkage software package implementing the CHP method. Since SEQLinkage can test for linkage heterogeneity and calculate Heterogeneity LOD (HLOD) scores the CHP method remains powerful when there is locus heterogeneity, i.e. the underlying genetic etiology is not due to the same gene/region in all families.

Materials and Methods

For the CHP method instead of analyzing each variant separately, multiple variants which form haplotypes within a genetic region, e.g. gene, are analyzed. This is done by constructing a marker which reflects the transmission pattern of the entire region and is numerically compatible with currently available linkage analysis methods and software. These markers incorporate allelic heterogeneity between and within families in a region and often have higher heterozygosity than SNVs, making them more informative and powerful to detect linkage.

To generate regional markers, haplotypes for the region must be obtained for all samples with sequence data. NGS data from family members are first checked for Mendelian errors and variants with Mendelian inconsistencies are removed. An improved version of the Lander-Green algorithm

for genetic phasing is applied to reconstruct haplotypes in the pedigrees⁵. For each pedigree, we first cluster variants on regional haplotypes by “bins”, e.g. LD blocks, and collapse variants in a bin into an indicator variable with values 0 or 1 for having no minor allele or at least one minor allele within the bin, which is similar to collapsing methods for rare variant association analysis⁶. We then assign each collapsed haplotype a single numeric value so that different patterns of collapsed haplotypes in each pedigree are uniquely represented (Figure 1). The choice of coding for patterns are arbitrary, although we use continuous positive integers and assign a smaller value for collapsed haplotypes having more 0’s than 1’s. The sample haplotypes thus represented can be directly used for parametric linkage analysis with many existing linkage software packages.

For WES data, genes can be used as regional markers. Within each region, commonly used bin size options for variants collapsing are 1) LD based collapsing, which uses estimated LD blocks as bins, 2) complete collapsing, whose bin size equals gene/region length and 3) no collapsing, whose bin size equals one. For regions where recombination events occur within a family, the sub-unit that shows the strongest evidence of linkage among all sub-units created by recombination breakpoints is used as the regional LOD score for the family, so that results from multiple families can still be combined.

In order to reconstruct genotypes for family members missing sequence data, linkage analysis requires marker allele frequencies. Frequencies of regional markers generated by CHP method can be derived from minor allele frequencies (MAF) of variants and pair-wise LD between variants. For rare variants with MAF derived from large samples (see Discussion), the minor allele counts can be approximated by a multivariate Poisson distribution with joint probability mass function

94 $P(\mathbf{X}) = f_{(\lambda, \theta)}(\mathbf{X})$ where $\lambda_{M \times 1}$ is expected allele counts for M variants and $\theta_{M \times M}$ is the variance-
 95 covariance matrix⁷. The covariance between variants X_i and X_j can be computed by
 96 $\text{cov}(x_i, x_j) = r_{ij} N \sqrt{p_i p_j (1 - p_i)(1 - p_j)}$ where r_{ij}^2 is the LD coefficient, p is population MAF
 97 and N is the sample size based on which population MAF are estimated. Therefore for a given
 98 haplotype pattern $\mathbf{x}_H = [x_1, x_2, \dots, x_M]$, $x_k \in \{0, 1\}$ the corresponding frequency $f_{(\lambda, \theta)}(\mathbf{X} = \mathbf{x}_H)$ can
 99 be computed from the probability mass function. When collapsing is applied, MAF for the
 100 collapsed unit is given as $1 - f_{(\lambda, \theta)}(\mathbf{X} = [0, 0, 0, \dots])$ by definition. Collapsed haplotype pattern
 101 frequencies thus computed are then used as the allele frequencies for the corresponding regional
 102 genotype markers.

103 To facilitate linkage analysis using sequence data in VCF format, we developed the SEQLinkage
 104 software that uses the Elston-Stewart algorithm as incorporated in FASTLINK⁸. It provides results
 105 in text format and high quality graphical reports for both LOD and HLOD scores. Additionally
 106 SEQLinkage supports output of regional genotype data into formats compatible with linkage
 107 software such as LINKAGE⁹ and Merlin¹⁰, with which two-point and multipoint parametric
 108 linkage analysis can be performed. Additionally MEGA2¹¹ format is supported, which can be used
 109 to transform data to the required input for a number of linkage programs.

110 To evaluate performance of our method we performed empirical type I error and power
 111 calculations for two-point linkage analysis using data on four non-syndromic hearing impairment
 112 (NSHI) genes: two autosomal recessive genes *GJB2* and *SLC26A4*, and two autosomal dominant
 113 genes *MYO7A* and *MYH9*. Two-generation pedigrees were simulated, with 3 to 8 offspring in the

114 last generation with the proportions determined by the distribution of number of children per
115 family in the United States in 2012, rescaled so that they sum to 100% (3 children: 69.34%, 4
116 children: 20.52%, 5 children: 6.84%, 6 children: 2.28%, 7 children 0.76%, 8 children 0.26%).
117 Genotypes are simulated for the four genes based on the variant sites and the corresponding minor
118 allele frequencies in European Americans recorded in the Exome Variant Server.

119 For type I error evaluations we use the same gene sequences and demographic data, yet simulate
120 disease pedigrees under the null, i.e., affection status not due to any of the rare variants in the gene
121 of interest. We consider different genetic architectures under the null including situations when 1)
122 variants in the gene region are in linkage equilibrium, 2) there is complete LD between variants
123 and 3) there exist within a gene recombination events in the sequence data of generated families.
124 Recombination events between variants are simulated based on rates obtained from Hapmap
125 Recombination Rates and Hotspots database (see Web Resources). Additionally we simulate
126 scenarios when parental genotypes are missing to evaluate type I error when CHP marker
127 frequencies have to be calculated using population MAF and LD estimated from data. Type I errors
128 are computed for cumulative HLOD scores on gene *SLC26A4* across 20 families using 2,000,000
129 replicates.

130 For power evaluations we annotate variants in these four NSHI genes using Deafness Variation
131 Database (DVD) and NCBI ClinVar, labelling variants as “causal” if they are so deemed by both
132 databases. Disease status for individuals are determined by genotypes on those causal sites under
133 dominant mode of inheritance for *MYO7A* and *MYH9*, and recessive (compound heterozygotes
134 and homozygotes) for *GJB2* and *SLC26A4*, assuming complete penetrance. Additionally for each

mode of inheritance we allow for allelic heterogeneity among families, i.e., the causal variant site in a gene may not be the same for different families. We “ascertain” simulated families having two or more affected offspring for linkage analysis. To introduce locus heterogeneity we sample families having causal variants in one gene but not the other, so that each simulated gene contributes to etiology of only a proportion of families in the entire dataset. We simulate 500 replicates under each different setting of sample size, mode of inheritance, presence of allelic heterogeneity and locus heterogeneity. For each replicate we compute LOD and HLOD scores using the CHP method. For comparison purposes we also analyze SNV markers and perform multipoint linkage analysis, with multipoint linkage analysis being performed using GeneHunter¹². Power is estimated by $P = \frac{N_{success}}{N}$ where the denominator is the total number of replicates and the numerator is the number of tests that successfully detected the linkage signal, i.e. LOD score greater than 3.3 or HLOD score greater than 3.6 which provides a genome wide significance level of $p < 0.05$ ¹³.

Results

Empirical type I error for the CHP linkage statistic is $\hat{\alpha} = 2.8 \times 10^{-5}$ (95% CI: $2.11 \times 10^{-5} \leq \alpha \leq 3.63 \times 10^{-5}$), demonstrating that type I error is well controlled and even conservative at a required α level of 4.7×10^{-5} for an HLOD of 3.6. Quantile-Quantile (QQ) plots are generated to evaluate the null distribution of test statistic in the presence of within-gene recombination, strong inter-marker LD and missing genotype data; type I error is well controlled and no sign of inflation is observed (Figure S1). Empirical power calculations for several known

non-syndromic hearing loss genes using the CHP method as well as for individual SNV are summarized by contour plots (Figures 2). Power analysis based on LOD and HLOD suggests that CHP is substantially more powerful for all models in the presence of intra- (Figure 2C) and inter-family allelic heterogeneity (Figures 2A–2C). For example to detect linkage with the *SLC26A4* gene using an autosomal recessive model with allelic heterogeneity, i.e. compound heterozygotes, and also with locus heterogeneity of 50%, 12 families are required for the CHP method to achieve a power of 90%, while analyzing individual SNVs requires >50 families to achieve the same power at a genome wide significance level of $\alpha=0.05$. Additionally, although multipoint linkage analysis is more powerful than analyzing SNVs, the CHP method is considerably more powerful than multipoint linkage analysis (Table S1).

For sequence data, variants are sometimes missing due to the inability to call variants or during quality control, variant calls are removed because of poor data quality. Therefore we also estimated sample size requirements for the CHP method when causal variants are missing from sequence data in a large proportion of families, i.e. 75%. The CHP method can tolerate missing data and is also always more powerful than the SNV method when there is missing data (Table 1).

Discussion

For linkage analysis, correct specification of marker allele frequency is crucial for controlling type I error and reducing type II error¹⁴. The number of founders with available genotypes in data for linkage analysis might often be too small to obtain a sufficiently accurate allele frequency estimate, thus we recommend the input VCF file be annotated with an external source of MAF information,

e.g. 1000 Genomes or Exome Variant Server. For some populations MAF information may not be available and frequencies estimated from founders have to be used.

In the context of Mendelian disease mapping it is reasonable to assume that common variants (variants having population $MAF > 1\%$) are not directly causal. Analyzing common variants will neither contribute to nor reduce power when analyzed with rare variants. Common variants can be in strong LD with variants in neighboring regions which may contain causal variants; thus when the CHP method is used to construct the regional marker also using common variants, linkage can be detected even though the region does not harbor any causal variants. Although common variants should not be used when constructing regional markers, we suggest analyzing common variants separately because they can potentially capture additional information when rare causal variants are missing from sequence data.

Analysis of rare variants using “burden” methods are usually limited to those variants which are most likely to be causal, e.g. missense, nonsense and splice site variants, because inclusion of non-causal variants can attenuate the association signal and reduce power. For the CHP methods inclusion of non-causal rare variants will not attenuate the linkage signal and therefore analysis does not need to be restricted to variants which are most likely functional and causal. Inclusion of non-causal rare variants to construct the region marker can provide additional linkage information if data for causal variants are missing. If the goal is to detect a linkage signal from variants which are potentially causal then linkage analysis using the CHP method can be limited to those variant sites which are most likely functional.

195 In addition to the CHP method being more powerful than performing multipoint linkage analysis,
196 it also controls type I error when there is missing parental genotype data and inter-marker LD,
197 which is not the case for multipoint linkage analysis. Caution should be used when performing
198 multipoint linkage analysis on sequence data, since when parental genotypes are missing for some
199 samples (common for NGS based family data) variants in LD can lead to serve inflated type I error
200 when markers are assumed to be in linkage equilibrium^{15,16}. The majority of multipoint linkage
201 analysis programs e.g. GeneHunter, SuperLink¹⁷, Vitesse¹⁸, do not take into consideration LD
202 between marker loci. Even for linkage programs that can model inter-marker LD, e.g.,
203 LINKAGE/FASTLINK and Merlin, the haplotype frequency estimates involving rare variants can
204 be inaccurate for studies with limited number of founders, leading to inflated type I error.

205 The SEQLinkage package, freely available at URL <http://bioinformatics.org/seqlink>, can
206 efficiently extract genotypes from VCF files and uses the CHP method described here to perform
207 linkage analysis as well as data format conversion on sequence data so that other programs can
208 also be used to perform linkage analysis if desired. It provides a novel and effective approach that
209 brings back well established linkage analysis techniques for use with the growing wealth of
210 genomic data of human pedigrees. Unlike filtering approaches which are commonly used to
211 analyze sequence data, SEQLinkage provides statistical evidence of the involvement of variants
212 in the etiology of Mendelian diseases. Additionally because it incorporates mode of inheritance
213 information and penetrance models it is less likely than filtering approaches to exclude causal
214 variants in the presence of phenocopies and/or reduced penetrance. For Mendelian traits for which
215 the penetrance model is not well established but the mode of inheritance is known, an affected-
216 only analysis can be performed where all unaffected individuals are made unknown to avoid

217 decreased power due the use of an incorrect penetrance model. We recommend the use of
218 SEQLinkage in parallel to filtering methods on the same sequence data to take full advantage of
219 the power of NGS in families.

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- 267

268 **Web Resources**

- 269 America's Families and Living Arrangements, <https://www.census.gov/prod/2013pubs/p20->
270 [570.pdf](https://www.census.gov/prod/2013pubs/p20-570.pdf)
- 271 Exome Variant Server (EVS), <http://evs.gs.washington.edu/EVS>
- 272 Deafness Variation Database (DVD), <http://deafnessvariationdatabase.com>

- 273 NCBI ClinVar, <https://www.ncbi.nlm.nih.gov/clinvar>
- 274 Hapmap Recombination Rates and Hotspots database,
- 275 <http://hapmap.ncbi.nlm.nih.gov/downloads/recombination/latest/rates/>

Figure legends

Figure 1. Coding of regional markers using the Collapsed Haplotype Pattern (CHP) method.

Three two-generational autosomal recessive pedigrees display the coding for a regional marker using information from six variant sites. Panel A shows two families segregating the same autosomal recessive disease which is due to different causal variants. Treating the entire region as a bin to collapse the variants effectively captures transmission of disease variants and allows for linkage information for a region to be summed across families. For regions with more diverse rare variant architecture as displayed in Panel B, where for this example disease etiology is caused compound heterozygotes variants, coding which represents both rare variant haplotypes is used to ensure that all meioses are informative. It should be noted that if coding as is shown in Panel A is used in this situation there will be a loss of information because all heterozygous offspring will be uninformative for linkage information, e.g. the meioses to offspring II:1 and II:4.

Figure 2. Power comparisons for LOD and HLOD statistics in two-point linkage analyses.

This figure shows the power for collapsed haplotype pattern markers (CHP) vs. single nucleotide variant (SNV) analysis under various modes of inheritance in the presence of intra- and inter-family allelic heterogeneity. X-axis is number of families, Y-axis is proportion of locus heterogeneity, i.e. the proportion of families with non-syndromic hearing impairment (NSHI) caused by detrimental variants in the gene under investigation, i.e. either *MYO7A* or *MYH9* for dominant model, or *GJB2* or *SLC26A4* for recessive model. Contour curves on the graphs are power estimates, dark orange lines for the CHP method and light blue lines for SNV analysis. Panel A displays the power for the LOD and HLOD statistics under an autosomal dominant model; panel B displays the power for the LOD and HLOD statistics under an autosomal recessive model; panel C displays the power for the LOD and HLOD statistics under an autosomal recessive model

299 in the presence of intra-family allelic heterogeneity, i.e. affected individuals are compound
300 heterozygous. CHP method is more powerful for both LOD and HLOD at a genome-wide
301 significance level of $\alpha=0.05$, but the absolute power of HLOD is not significantly larger than LOD.
302 This is due to the very low MAFs for the genes under study and therefore for most families all
303 variants in the non-causal gene are monomorphic and therefore are uninformative.

304

306 **Table 1: Sample size estimates for the simulated nonsyndromic hearing impairment study.**

307

Required Power	Gene	MOI	CHP ¹	SNV ²	CHP-M75% ³	SNV-M75%
0.8	<i>SLC26A4</i>	recessive	11	40	39	160
0.9	<i>SLC26A4</i>	recessive	13	45	46	180
0.8	<i>SLC26A4</i>	compound recessive	11	50	39	200
0.9	<i>SLC26A4</i>	compound recessive	13	55	46	220
0.8	<i>GJB2</i>	recessive	12	23	44	92
0.9	<i>GJB2</i>	recessive	14	28	52	112
0.8	<i>GJB2</i>	compound recessive	12	25	44	100
0.9	<i>GJB2</i>	compound recessive	14	34	52	136
0.8	<i>MYO7A</i>	dominant	12	16	31	64
0.9	<i>MYO7A</i>	dominant	14	20	36	80
0.8	<i>MYH9</i>	dominant	11	13	32	52
0.9	<i>MYH9</i>	dominant	14	18	41	72

Note: 50% locus heterogeneity is assumed for all scenarios.

¹Number of families required for CHP method.

²Number of families required for single variant method.

³“M-75%”: number of families required when causal variants in 75% participating families are missing.