GENOMIC REGIONS LINKED TO SOFT SWEEPS APPROXIMATE NEUTRALITY WHEN INFERRING POPULATION HISTORY FROM SITE PATTERN FREQUENCIES

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ABSTRACT

Recent studies have suggested that selection is widespread throughout the genome and largely uncompensated for in inferences of population history. To address this potential issue, we estimated site pattern frequencies for neutral and selection associated areas of the genome. There are notable differences in these frequencies between neutral regions and those affected by selection. However, these differences have relatively small effects when inferring population history.

1 INTRODUCTION AND BACKGROUND

In the past year, population geneticists have been debating the extent to which natural selection has shaped the human genome. Evidence suggests that soft sweeps (Harris et al., 2018; Schrider and Kern, 2017) and polygenic adaptation (Daub et al., 2013; Hernandez et al., 2011; Pritchard et al., 2010) are the primary modes of selection in humans. This has led some researchers to suggest that most of the genome is in some way affected by selection either directly or indirectly through linkage with neighboring sites. This led Kern and Hahn (2018) to argue that the original lines of evidence that led to the neutral theory of evolution pioneered by Kimura (1983) do not hold up in the genomic era. Specifically, they believe inferred population histories are skewed because areas of the genome often perceived to be neutral are actually affected by selection. In addition, they suggest genome-wide selection scans will have a high rate of false negatives if using a null distribution built from selected regions of the genome. On the other hand, Harris et al. (2018) and Jensen et al. (2018) suggest that many of these apparent signals of selection are false positives, and

others have suggested that selection has been less common but signals of selection are amplified by population history (Torres et al., 2018).

A simple way to test the idea proposed by Kern and Hahn (2018) is to reconstruct population history using different areas of the genomes. Schrider and Kern (2017) used a machine learning algorithm to assemble a list of regions in the genome inferred to be affected by selection but previously thought to be neutral, and those that are neutral. Here, relative site pattern frequencies (Rogers, 2019) are calculated in each of these subdivisions to measure the potential effects of selection. In what follows, we show that selection skews site pattern frequencies, but has little effect on the estimation of population history in our model.

2 RESULTS

Selection affects population history. A nucleotide site pattern is a particular arrangement of derived and ancestral alleles when a single haploid individual is sampled from each sample population. The total number of distinct site patterns is therefore all combinations in which at least one sample, but not all, carries the derived allele. Three populations are discussed here, given the labels CEU (X), JPT (Y), and YRI (Z). The possible site patterns are x , y , z , xy , xz , and yz , the first three representing the cases in which the derived allele is found in only one population, and the rest representing when the derived allele is found in two populations. Estimations of common ancestry, divergence times, and admixture can be made using the relative frequency of these site patterns (Rogers, 2019). The difference between the selection-affected site patterns and neutral site patterns was calculated. If neutral and selection-affected genomic regions have similar site pattern frequencies to neutral regions, it will produce similar estimates of population history, and the difference between them should not vary from zero significantly.

Figure 1 shows the difference in relative frequencies of site patterns between affected and neutral regions for biallelic single nucleotide polymorphisms (SNP). Site pattern frequencies were calculated using the *sitpat* program from the *Legofit* package (Rogers, 2019). Confidence intervals were generated by using 1,000 bootstrap replicates generated from *sitpat*. Neutral and selection-affected regions differed significantly in the x , y , and xy site patterns, with confidence intervals that do not overlap with zero. To investigate the possibility that this pattern is driven primarily by one type of selection, hard sweeps or soft sweeps, affected regions were split accordingly. Hard sweeps differ from other distinctions substantially but the confidence intervals are large, likely due to the relatively small sample size of these regions (Table 1). Soft sweep regions are marginally more similar to neutral regions, but the large difference between affected and neutral regions in the x , y , and xy patterns persists when only soft sweeps are considered.

Inference of Population History. Site pattern frequencies differ between neutral and selected affected regions of the genome. However, these differ-

ences are small and it is unclear how large their effect will be when inferring population history. To test the effect these differences have, the site patterns for neutral and soft-sweep affected regions were used to estimate ancestral divergence times and population sizes using *Legofit* (Rogers, 2019). The demographic model used was taken from (Rogers et al., 2019). This model includes admixture events from Neanderthals into Eurasia, ancient humans into Neanderthals, and from superarchaic hominins into Denisovans and the ancestor of Neanderthals and Denisovans. Rogers et al. (2019) found that the exclusion of these admixture events can strongly bias results.

Selection seems to affect the estimation of these parameters, but only to a

Figure 1: Difference in selection-affected and neutral relative site pattern frequencies. If all patterns rested at zero, the inferred population history would be identical. Each population is given an alphabetical label: x (CEU), y (JPT), z (YRI)

Samples	Neutral	Selection Affected	Soft-Sweep Affected	Hard-Sweep Affected
Humans only	285,833	619,136	431,105	4,460
Human and Neanderthal	251,953	543,098	378,700	4,104
Human, Neanderthal, and Denisova	261,996	563,523	392,625	4,275

Table 1: Number of sites tabulated in each run of *sitepat*

85 small extent. Figure 2 shows the percent differences between soft-affected and
86 neutral estimates of divergence times. Confidence intervals were generated by
87 taking differences of individual estimates in each of fifty bootstrap replicates.
88 If selection does not strongly affect quantitative estimates of demographic in-
89 ference, these differences should rest around zero.

Figure 2: Percent difference between estimates of divergence times for soft-sweep associated regions and neutral regions for a model including Yorubans (x), European (CEU) (y), Neanderthals (n) further split into the Vindija (v) and Altai (a), Denisovans (d), and superarchaic hominins (s).

90 Figure 3 shows the percent differences between soft-affected and neutral es-
91 timates of ancestral population sizes. In general, estimates of population size
92 were more likely to be disrupted by the effects of soft selective sweeps. How-

93 ever, these deviations are relatively small. The largest difference in population
94 size without confidence intervals overlapping with zero is approximately one
95 or two percent of approximately 40,000 (Table 2).

Table 2: Estimates of divergence times and population sizes for a model including Yorubans (x), Europeans (CEU) (y), Neanderthals (n), Denisovans (d), and superarchaic hominins (s). “T” indicates divergence time and “2N” is the diploid effective population size.

Parameter	Neutral	Soft Selection Affected
T_{xynds}	67,049.1	52,396.2
T_{nd}	24,906.3	24,921.1
T_{av}	15,570.6	15,617.5
T_{xy}	3,864.6	3,628.0
T_d	1,855.0	1,906.8
T_a	4,958.1	4,883.7
T_v	2,339.7	2,315.4
$2N_{av}$	31,673.6	24,333.9
$2N_n$	7,372.7	7,414.5
$2N_{nd}$	2,111.9	2,063.7
$2N_{xy}$	50,259.1	50,639.3
$2N_{xynd}$	42,152.5	42,826.4
$2N_s$	81,794.2	50,639.7

96 3 DISCUSSION

97 The difference in site pattern frequencies is in agreement with the results of
98 Schrider and Kern (2017). The site patterns for the neutral regions imply a
99 different population history than the selection affected regions. The neutral
100 and selection-affected regions are close for each site pattern except the x and
101 xy patterns. The x and y site patterns are relatively over-represented in the
102 neutral case, while the xy site pattern is relatively underrepresented. The se-
103 lection affected site patterns would therefore overestimate the length of the
104 ancestral Eurasian branch, and underestimate the individual Eurasian popula-
105 tion branches.

106 The significant differences in site pattern frequencies do not translate to
107 large differences in estimations of population history in our model. In the Eu-
108 ropean model, only one estimated parameter, the size of the ancestral popula-
109 tion of humans, Neanderthals, and Denisovans ($2N_{xynd}$), shows a significant
110 difference between neutral and soft sweep affected regions. However, in this
111 case the difference is more than an order of magnitude smaller than the esti-
112 mated parameter itself.

113 The results here are consistent with large portions of the genome being un-

Figure 3: Percent difference between estimates of ancestral effective population size for soft-sweep associated regions and neutral regions for a model including Yorubans (x), European (CEU) (y), Neanderthals (n) further split into the Vindija (v) and Altai (a), Denisovans (d), and superarchaic hominins (s).

Figure 4: Percent difference shown in 3 zoomed in to compensate for the visual bias caused by the large amount of error around estimates of $2N_{xynd}$.

der selection. However, differences in site pattern frequencies generated by soft selection do not appear to have substantial effects on the estimation of parameters of population history, at least when using *Legofit*. This result may have a larger effect at finer scales. For instance, the significant difference in T_{av} , the divergence time of the Altai and Vindija Neanderthals, could be as large as 5,000 generations or 125,000 years. This may not too large when considering recent human population history and admixture between humans and other hominins, but could be problematic in reconstructing a fine scale history of Neanderthal populations. Further work exploring different time scales and population histories should be done before any generalization of these results is made. Nonetheless, soft sweeps do not cause meaningful disturbances in estimates of population history at the scale studied here.

4 METHODS

Simons Data. Simons Genome Diversity Mallick et al. (2016) data for Japan (JPT), Yoruba (YRI), and Europe (England and France) was acquired from <https://www.ebi.ac.uk/ena/data/view/PRJEB9586>. Sites with a map quality or genotype quality below thirty were excluded. Sites that were fixed across all human populations were removed, because they cannot differentiate human populations. Indels, SNPs within seven bases of an indel, and low quality SNPs (filter level equal to zero) were removed. The Central Europeans from Utah (CEU) sample is not represented in the SGDP. Instead, individuals from England and France were used as a proxy for CEU. These samples were used because CEU is present in the analysis and results of Schrider and Kern (2017), but absent from the Simons data.

Subdivisions. Schrider and Kern (2017) subdivide the genome into neutral and selection linked, soft-sweep affected, and hard-sweep affected regions. These results were obtained from <https://github.com/kern-lab/shIC/tree/master/humanScanResults>. The selection linked regions are those that are commonly assumed to be neutral in the literature, but showed evidence of being linked to and affected by selection in their machine learning analysis. The selection-affected and neutral regions were used here to divide data respectively. Regions inferred to be under selection are not studied here because selected regions are already expected to produce different estimates of population history from neutral regions. Here we are concerned with regions of the genome that could be mistaken for neutral regions and skew population histories in the literature. Each sample has its own subdivision that reflects the population and selective history of the population it belongs to. The neutral distinction made here refers to sites where all three populations are considered neutral. For a site to be included in the “selection-affected” for the purpose of this analysis, at least one population needs to be represented in the original “selection-affected” distinction.

Site Pattern Frequencies. The program *sitapat* from the *Legofit* (Rogers, 2019) package was used to calculate relative site pattern frequencies (SPF) (Rogers

et al., 2017). The ancestral allele is determined by using reference alleles of chimpanzees and gorillas. The analysis was limited to sites where the chimpanzee and gorilla were fixed for the same allele, and the human samples were polymorphic. One thousand bootstrap replicates were generated for each combination of selection type and set of populations. The difference in site pattern frequencies between selection affected regions and neutral regions was then taken, with confidence intervals generated from differences in individual bootstrap replicates.

Legofit. Site pattern frequencies were used to estimate population history parameters using *Legofit*. The model of population history includes admixture between Eurasians and Neandethals, and between superarchaics and Denisovans and the ancestor of Denisovans and Neanderthals. Site pattern frequencies are generated using high coverage Neanderthal genomes from the Vindija (Prüfer et al., 2017) and Altai (Prüfer et al., 2014) Neanderthals, and a high coverage Denisovan genome from Siberia (Meyer et al., 2012). The Yoruban samples from the SGDP represent Africa. The final human population was a mixture of English and French individuals serving as a proxy for CEU. One thousand bootstrap replicates were generated for each model of population history. Confidence intervals for these differences were generated by taking the inner ninety-five percent of the differences between individual bootstrap replicate estimations. This process was conducted for neutral and soft-selection affected regions separately.

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