

Discovering epistasis between germline mutator alleles in mice

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Abstract

Maintaining genome integrity in the mammalian germline is essential and enormously complex. Hundreds of proteins comprise pathways involved in DNA replication, and hundreds more are mobilized to repair DNA damage [1]. While loss-of-function mutations in any of the genes encoding these proteins might lead to elevated mutation rates, *mutator alleles* have largely eluded detection in mammals.

DNA replication and repair proteins often recognize particular sequence motifs or excise lesions at specific nucleotides. Thus, we might expect that the spectrum of *de novo* mutations — that is, the frequency of each individual mutation type (C>T, A>G, etc.) — will differ between genomes that harbor either a mutator or wild-type allele at a given locus. Previously, we used quantitative trait locus mapping to discover a mutator allele near the DNA repair gene *Mutyh* that increases the rate of *de novo* C>A germline mutation in a collection of recombinant inbred lines (RILs) known as the BXDs [2,3].

In this study, we developed a new method to detect alleles that affect the mutation spectrum in biparental RILs. By applying this approach to mutation data from the BXDs, we confirmed the activity of the germline mutator locus near *Mutyh* and discovered an additional C>A germline mutator locus on chromosome 6 that overlaps *Ogg1* and *Mbd4*, two DNA glycosylases involved in base-excision repair [4,5]. Strikingly, BXDs with mutator alleles on chromosome 6 only exhibit elevated rates of C>A germline mutation if they also possess mutator alleles near *Mutyh*, and BXDs with both alleles exhibit even higher C>A mutation rates than those with either one alone.

To our knowledge, these new methods for analyzing mutation spectra reveal the first evidence of epistasis between mammalian germline mutator alleles, and may be applicable to mutation data from humans and other model organisms.

Introduction

The germline mutation rate is a fundamental parameter in population genetics, and reflects the complex interplay between DNA replication and repair pathways, exogenous sources of DNA damage, and life-history traits. For example, parental age is an important determinant of mutation rate variability; in many mammalian species, the number of germline *de novo* mutations observed in offspring increases as a function of paternal and maternal age [6,7,8,9,10]. Rates of germline mutation accumulation are also variable across human families [7,11], potentially due to genetic variation or differences in environmental exposures. Nonetheless, genetic variants that augment germline mutation rates, known as *mutator alleles*, have largely eluded detection in mammals.

Studies of thousands of human trios have failed to identify common variants associated with germline mutation rate variation [12], meaning that any mutators that segregate within the human population are likely rare or modest in effect size. The dearth of observed germline mutators in mammalian genomes is not terribly surprising, since alleles that lead to elevated germline mutation rates would likely have deleterious consequences and be purged by negative selection [13]. Moreover, germline mutation rates are relatively low, and direct mutation rate measurements generally require whole-genome sequencing data from both parents and their offspring. As a result, large-scale association studies — which have been used to map genetic contributions to numerous other complex traits — are not well-powered to investigate the polygenic architecture of germline mutation rates.

Despite these challenges, less traditional strategies have been successfully used to identify a small number of mutator alleles in both humans and mice. By focusing on children who suffer from rare

genetic diseases, a recent study identified two mutator alleles that lead to significantly elevated rates of *de novo* germline mutation in human families [14]. Another group discovered mutator phenotypes in the sperm and somatic tissues of adults that suffer from cancers caused by inherited mutations in the POLE/POLD1 exonucleases [15].

In mice, candidate mutator alleles have also been discovered by sequencing a large collection of recombinant inbred lines (RILs). Commonly known as the BXDs [3], these RILs were derived from either F2 or advanced intercrosses of C57BL/6J and DBA/2J, two laboratory strains that exhibit significant differences in their germline mutation spectra [16]. The BXD RILs were maintained via brother-sister mating for up to 180 generations, and each line therefore accumulated hundreds or thousands of germline mutations on a nearly-homozygous linear mosaic of parental B and D haplotypes. Due to their husbandry in a controlled laboratory setting, the BXDs were largely free from confounding by environmental heterogeneity, and the effects of selection on *de novo* mutations were attenuated by strict inbreeding [17].

Whole-genome sequencing data from 152 BXD strains were previously used to map a quantitative trait locus (QTL) for the C>A germline *de novo* mutation rate. Germline C>A mutation rates were nearly 50% higher in mice with D alleles at the QTL, likely due to the reduced efficacy of oxidative DNA damage repair caused by genetic variation in the DNA glycosylase *Mutyh*. Importantly, the quantitative trait locus was undetectable in a genome-wide scan for variation in overall mutation rates, which were only modestly higher in BXDs with D alleles, demonstrating the utility of *mutation spectrum* analysis for mutator allele discovery. Close examination of the mutation spectrum is likely to be broadly useful for detecting mutator alleles, as genes involved in DNA replication and repair often recognize particular sequence motifs or excise specific types of DNA lesions [18]. Mutation spectra are usually defined in terms of k -mer nucleotide context [18]; the 1-mer mutation spectrum, for example, consists of 6 mutation types after collapsing by strand complement (C>T, C>A, C>G, A>T, A>C, A>G), while the 3-mer mutation spectrum contains 96 (each of the 1-mer mutations partitioned by trinucleotide context).

Although mutation spectrum analysis can enable the discovery of mutator alleles that affect the rates of specific mutation types, early implementations of this strategy have suffered from some drawbacks. For example, performing association tests on the rates or fractions of every k -mer mutation type can quickly incur a substantial multiple testing burden. Since germline mutation rates are generally quite low, estimates of k -mer mutation type frequencies from individual samples may also be noisy and imprecise. We were therefore motivated to develop a statistical method that could overcome the sparsity of *de novo* mutation spectra, eliminate the need to test each k -mer mutation type separately, and enable sensitive detection of alleles that influence the germline mutation spectrum.

Here, we present a new mutation spectrum association test that minimizes multiple testing burden and mitigates the challenges of sparsity in *de novo* mutation datasets. We leverage this method to re-analyze germline mutation data from the BXD cohort, and find compelling evidence for the existence of a second mutator allele that was undetectable using previous approaches. The new allele appears to interact epistatically with the mutator that was previously discovered in the BXDs, further augmenting the C>A germline mutation rate in a subset of RILs. Our observation of epistasis suggests that mild DNA repair deficiencies may compound one another, as mutator alleles chip away at the redundant DNA repair systems that collectively maintain germline integrity.

Results

A novel method for detecting mutator alleles

We developed a statistical method, termed “inter-haplotype distance” (IHD), to detect loci that are associated with mutation spectrum variation in biparental RILs (Figure 1). Our approach leverages the fact that mutator alleles often leave behind distinct and detectable impressions on the *mutation spectrum*, even if they increase the overall mutation rate by a relatively small amount (Materials and Methods). Given a population of haplotypes, we assume that a) each has been genotyped at the same collection of biallelic loci and b) each harbors *de novo* mutations that have been partitioned by k -mer context (Figure 1). At every locus, we calculate a cosine distance between the aggregate mutation spectra of haplotypes that inherited either allele. Using permutation tests, we then identify loci at which those distances are larger than what we’d expect by random chance.

We find that the method’s power is primarily limited by the initial mutation rate of the k -mer mutation type affected by a mutator allele and the total number of *de novo* mutations used to detect it (Figure 1-[figure supplement 1](#)). For example, given 100 haplotypes with an average of 500 *de novo* germline mutations each, the IHD approach has approximately 80% power to detect a mutator allele that increases the C>T *de novo* mutation rate by as little as 10%. However, the method only has about 10% power to detect a mutator of identical effect size that augments the C>A mutation rate, since C>A mutations are expected to make up a smaller fraction of all *de novo* germline mutations to begin with. These simulations also demonstrate that our approach is well-powered to detect large-effect mutator alleles (e.g., those that increase the mutation rate of a specific k -mer by 50%), even with a relatively small number of mutations per haplotype (Figure 1-[figure supplement 1](#)). Overall, we find that both IHD and traditional quantitative trait locus (QTL) mapping have similar power to detect alleles that augment the rates of individual 1-mer mutation types (Figure 1-[figure supplement 2](#)); for a more detailed discussion of the advantages and disadvantages of either method, see the *Supplementary Information*.

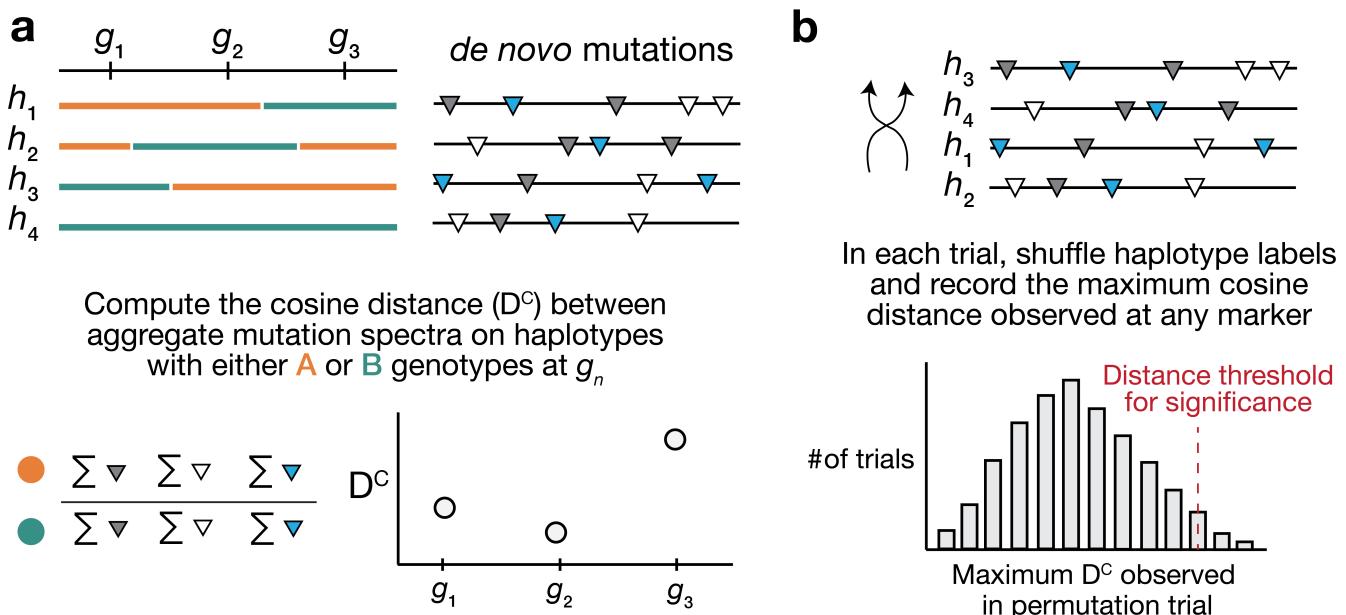


Figure 1: Overview of inter-haplotype distance method for discovering mutator alleles. **a)** A population of four haplotypes has been genotyped at three informative markers (g_1 through g_3); each haplotype also harbors private *de novo* germline mutations, which are denoted as triangles. *De novo* mutations are further classified into three possible mutation types, indicated by the color of each triangle. At each informative marker g_n , we calculate the total number of each mutation type observed on haplotypes that carry either parental allele (i.e., the aggregate mutation spectrum). We then calculate the cosine distance between the two aggregate mutation spectra. We repeat this process for every informative marker g_n . **b)** To assess the significance of any distance peaks in a), we perform permutation tests. In each of N permutations, we shuffle the haplotype labels associated with the *de novo* mutation data, run a genome-wide distance scan, and record the maximum cosine distance encountered at any locus in the scan. Finally, we calculate the $1 - p$ percentile of the distribution of those maximum distances to obtain a genome-wide cosine distance threshold at the specified value of p .

Re-identifying the mutator allele on chromosome 4 in the BXDs

We applied our inter-haplotype distance method to 93 BXD RILs (Materials and Methods) with a total of 62,993 *de novo* germline mutations [2]. Using mutation data that were partitioned by 1-mer nucleotide context, we discovered a locus on chromosome 4 that was significantly associated with mutation spectrum variation (Figure 2A; maximum adjusted cosine distance of 352.7 at marker ID rs52263933 ; position 116.75 Mbp in GRCm38/mm10 coordinates).

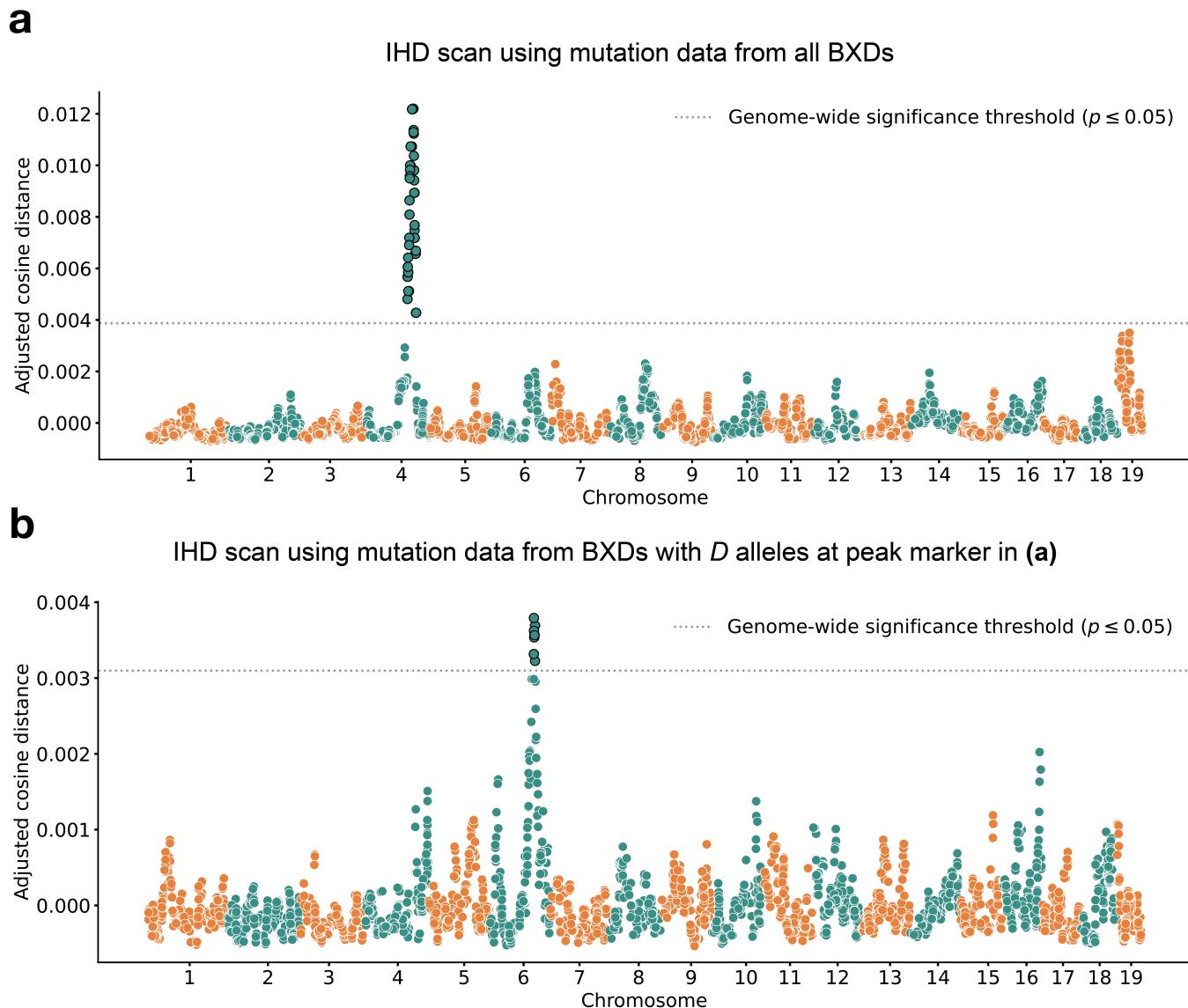


Figure 2: Results of inter-haplotype distance scans in the BXD RILs. **a)** Adjusted cosine distances between aggregate 1-mer *de novo* mutation spectra on BXD haplotypes ($n = 93$ haplotypes; 62,993 total mutations) with either *D* or *B* alleles at 7,320 informative markers. Cosine distance threshold at $p = 0.05$ was calculated by performing 10,000 permutations of the BXD haplotype mutation data, and is shown as a dotted grey line. **b)** Adjusted cosine distances between aggregate 1-mer *de novo* mutation spectra on BXD haplotypes with *D* alleles at rs52263933 ($n = 55$ haplotypes; 40,913 total mutations) and either *D* or *B* alleles at 7,320 informative markers. Cosine distance threshold at $p = 0.05$ was calculated by performing 10,000 permutations of the BXD haplotype mutation data, and is shown as a dotted grey line.

In a previous analysis, we used quantitative trait locus (QTL) mapping to identify a nearly identical locus on chromosome 4 that was significantly associated with the C>A germline mutation rate in the BXDs [2]. This locus overlapped 21 protein-coding genes that are annotated by the Gene Ontology as being involved in “DNA repair,” but only one of these genes contained non-synonymous differences between the two parental strains: *Mutyh*. *Mutyh* encodes a protein involved in the base-excision

repair of 8-oxoguanine (8-oxoG), a DNA lesion caused by oxidative damage, and prevents the accumulation of C>A mutations [4,19,20]. C>A germline mutation rates are nearly 50% higher in BXDs that inherit *D* genotypes at marker ID rs52263933 (the marker at which we observed the highest adjusted cosine distance on chromosome 4) than in those that inherit *B* genotypes (Figure 3) [2].

An additional germline mutator allele on chromosome 6

After confirming that the inter-haplotype distance method could recover the mutator locus overlapping *Mutyh*, we asked if our approach could identify additional mutator loci in the BXD. To account for the effects of the large-effect mutator locus near *Mutyh*, we divided the BXD RILs into those with either *D* (n = 55) or *B* (n = 38) genotypes at rs52263933 and ran a genome-wide distance scan using each group separately (Figure 2B).

Using the BXDs with *D* genotypes at rs52263933, we identified a locus on chromosome 6 that was significantly associated with mutation spectrum variation (Figure 2B; maximum adjusted cosine distance of 81.0 at marker rs31001331; position 114.05 Mbp in GRCm38/mm10 coordinates). We performed QTL scans for the fractions of each 1-mer mutation type using these same mutation data, but none produced a genome-wide significant log-odds score at any locus (Figure 2-figure supplement 1; Materials and Method). We queried the region surrounding the locus on chromosome 6 (+/- 5 Mbp) and discovered 15 protein-coding genes that harbored nonsynonymous differences between the parental C57BL/6J and DBA/2J strains. Two of these genes were also annotated with the Gene Ontology term “DNA repair”: *Ogg1* and *Mbd4*. *Ogg1* encodes a key member of the base-excision repair response to oxidative DNA damage (a pathway that also includes *Mutyh*), and *Mbd4* encodes a protein that is involved in the repair of G:T mismatches at methylated CpG sites that have undergone spontaneous deamination. Each of these genes harbors a single fixed nonsynonymous difference between the C57BL/6J and DBA/2J parental strains (Table 1).

Table 1: Nonsynonymous mutations in DNA repair genes near the chr6 peak

Gene name	Ensembl transcript name	Nucleotide change	Amino acid change	Position in GRCm38/mm10 coordinates	SIFT prediction
<i>Ogg1</i>	ENSMUST00000032406	A>G	p.Thr95Ala	chr6:113,328,510	0.84 (tolerant/benign)
<i>Mbd4</i>	ENSMUST00000032469	C>T	p.Asp129Asn	chr6:115,849,644	0.02 (intolerant/delete rious)

We also considered the possibility that expression quantitative trait loci (eQTLs), rather than nonsynonymous mutations, could contribute to the C>A mutator phenotype associated with the locus on chromosome 6. Using GeneNetwork [21] we mapped cis-eQTLs for *Ogg1* and *Mbd4* in a number of tissues, including hematopoietic stem cells, kidney, and spleen; we did not have access to expression data from germline tissues. BXD genotypes near the cosine distance peak on chromosome 6 were significantly associated with *Ogg1* expression in some (but not all) tissues, and *D* genotypes were nearly always associated with decreased gene expression (Table supplement 1). We discovered one significant cis-eQTL for *Mbd4* in spleen at which *D* alleles were associated with increased expression. We also queried a previously published collection of eQTLs derived from Diversity Outbred (DO) mouse embryonic stem cell expression data [22], but did not find any significant eQTLs for either *Ogg1* or *Mbd4*.

Finally, we queried the structural variants identified in a population of inbred laboratory strains sequenced by the Sanger Mouse Genomes Project (MGP) [23] and found 93 deletions within the implicated interval on chromosome 6. Of these, five overlapped the exonic sequences of three

protein-coding genes: *Tmem72*, *Tmcc1*, and *Gm20387*. None of these genes has a previously annotated role in DNA binding, repair or replication, or in a pathway that would likely affect germline mutation rates.

Evidence of epistasis between germline mutator alleles

Next, we more precisely characterized the effects of the chromosome 4 and 6 mutator alleles on mutation spectra in the BXDs. We observed that C>A germline mutation fractions in BXDs with *D* alleles at both mutator loci were higher than in BXDs with *D* alleles at either locus alone (Figure 3 and [2-figure supplement 2](#)). Compared to BXDs with *B* alleles at the chromosome 6 mutator locus, those with *D* alleles did not exhibit higher C>A mutation fractions, indicating that the effects of the chromosome 6 mutator locus depend on the presence of a *D* allele at the chromosome 4 locus (Figure 3).

We also used SigProfilerExtractor [24] to assign the germline mutations in each BXD to a single-base substitution (SBS) mutation signature from the COSMIC catalog [25]. Mutation signatures often reflect specific exogenous or endogenous sources of DNA damage, and the fraction of a given sample's mutations attributable to a particular SBS signature can suggest a genetic or environmental etiology. The SBS1, SBS5, and SBS30 mutation signatures were active in all BXDs, regardless of genotypes at the chromosome 4 and 6 mutator loci (Figure 3). However, the SBS18 signature, which is dominated by C>A mutations and likely reflects unrepaired DNA damage from reactive oxygen species, was only active in mice with *D* alleles at the chromosome 4 locus; the highest SBS18 activity was observed in mice with *D* alleles at both mutator loci (Figure 3). SBS18 activity was absent in mice with *D* alleles at the chromosome 6 mutator locus alone (Figure 3), further demonstrating that *D* alleles at this locus are not sufficient to cause a mutator phenotype.

To more formally test for epistasis, we fit a linear model predicting counts of C>A mutations in each strain as a function of genotypes at `rs52263933` and `rs31001331` (the peak markers at the two mutator loci) (*Materials and Methods*). A model that included an interaction term between genotypes at the two markers fit the data significantly better than a model including only additive effects ($p = 9.8e-4$).

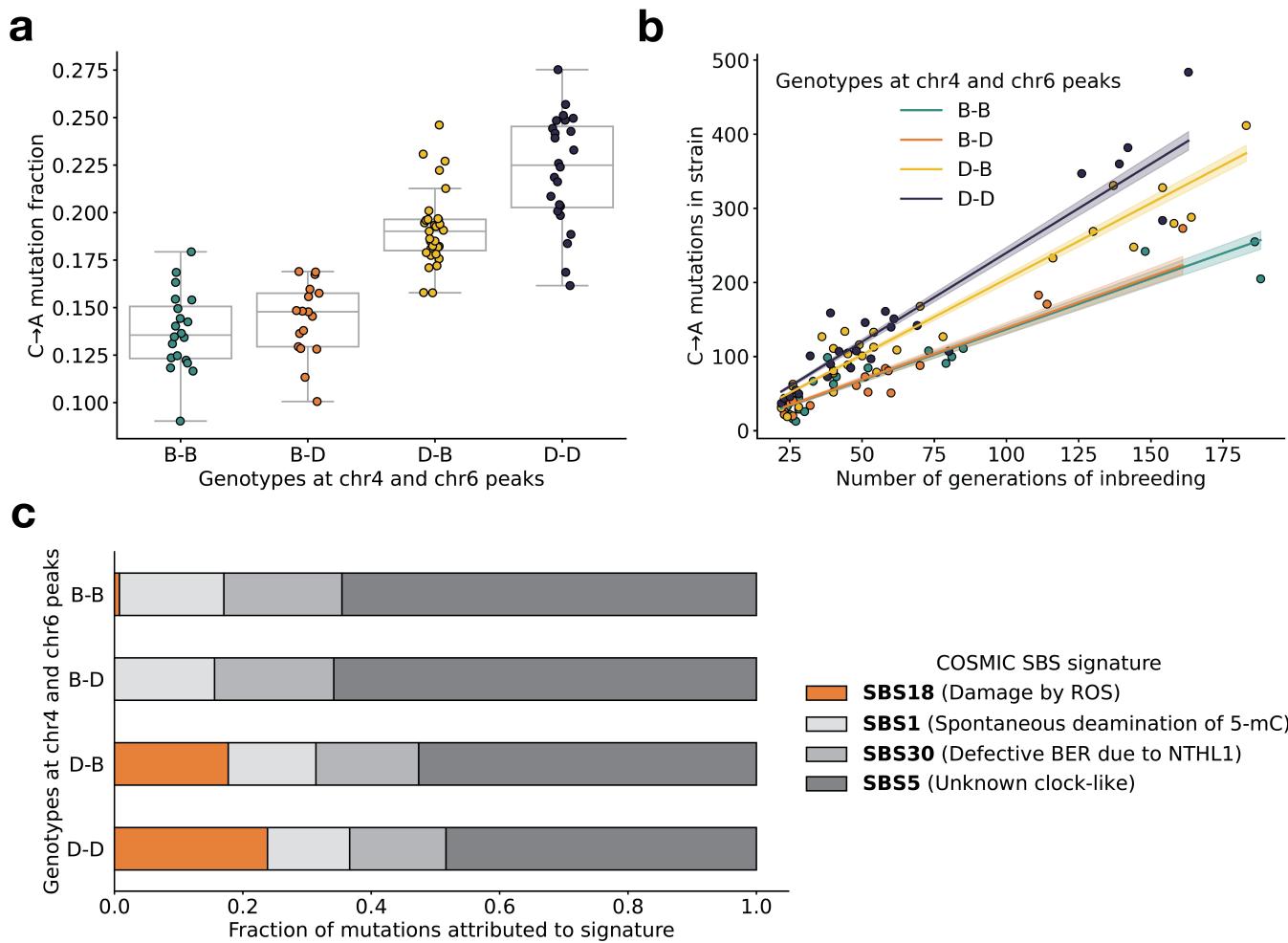


Figure 3: BXD mutation spectra are affected by alleles at both mutator loci. **a)** C>A *de novo* germline mutation fractions in BXDs with either *D* or *B* genotypes at markers rs52263933 (chr4 peak) and rs31001331 (chr6 peak). **b)** Counts of C>A *de novo* germline mutations in each BXD strain plotted against the number of generations for which it was inbred. Lines represent predicted C>A counts in each haplotype group from a Poisson regression (identity link), and shading around each line represents the 95% confidence interval for the predicted values. **c)** Germline mutations in each BXD were assigned to COSMIC SBS mutation signatures using SigProfilerExtractor. After grouping BXDs by their genotypes at rs52263933 and rs31001331, we calculated the fraction of mutations in each group that were assigned to each signature. The proposed etiologies of each mutation signature are: SBS1 (spontaneous deamination of methylated cytosine nucleotides) SBS5 (unknown, clock-like signature), SBS18 (damage by reactive oxygen species, possibly loss-of-function mutations in MUTYH), and SBS30 (defective base-excision repair due to NTHL1 mutations).

To explore the effects of the two mutator loci in other inbred laboratory mice, we also compared the germline mutation spectra of Sanger Mouse Genomes Project (MGP) strains [23]. Dumont [16] previously identified germline mutations that were private to each of the 29 MGP strains; these private variants likely represent recent *de novo* germline mutations (Figure 3—figure supplement 1). Only two of the MGP strains possess *D* genotypes at both the chromosome 4 and chromosome 6 mutator loci: DBA/1J and DBA/2J. As before, we tested for epistasis in the MGP strains by fitting two linear models predicting C>A mutation counts as a function of genotypes at rs52263933 and rs31001331. A model incorporating an interaction term between genotypes at these loci did not fit the MGP data significantly better than a model with additive effects alone ($p = 0.174$). Thus, we are unable to confirm the signal of epistasis observed in the BXDs, but this may be due to the smaller number of MGP strains with *de novo* germline mutation data.

Only one of the candidate mutator alleles is present in wild mice

To determine whether the candidate mutator alleles on chromosome 6 were segregating in natural populations, we queried previously published sequencing data generated from 67 wild-derived mice

[26]. These data include three subspecies of *Mus musculus*, as well as the outgroup *Mus spretus*. We found that the *D* allele of *Ogg1* was segregating at an allele frequency of 0.259 in *Mus musculus domesticus*, the species from which C57BL/6J and DBA/2J derive the majority of their genomes [27], and was fixed in *Mus musculus musculus*, *Mus musculus castaneus*, and the outgroup *Mus spretus*. However, the *D* allele of *Mbd4* was not present in any of the wild mice.

Discussion

Epistasis between germline mutator alleles

To our knowledge, these results reveal evidence of epistasis between mammalian germline mutator alleles for the first time. BXDs with *D* alleles at both the previously-identified mutator locus on chromosome 4 [2] and the novel locus on chromosome 6 have significantly higher C>A germline mutation fractions than lines with *D* alleles at either locus alone (Figure 3). Moreover, those with *D* alleles at the chromosome 6 locus don't exhibit elevated C>A mutation rates, suggesting that those alleles are not sufficient to cause a mutator phenotype. Our observations in the BXDs raise the exciting possibility that epistasis between mutator alleles has contributed to the evolution of germline mutation rates and spectra in mammalian genomes.

Importantly, we discovered evidence of epistasis between germline mutator alleles in an unnatural population; the BXDs were inbred by brother-sister mating in a highly controlled laboratory environment that attenuated the effects of natural selection on all but the most deleterious alleles [17]. Large-effect mutator alleles (and epistasis between them) may be much less common in natural, outbreeding mammalian populations. Regardless, our results demonstrate that germline mutation rates in recombinant inbred populations are highly plastic, and that RILs represent a powerful system in which to discover germline mutators.

Potential causal variants underlying the mutator allele

Two protein-coding DNA repair genes overlap the C>A mutator locus on chromosome 6 and also contain nonsynonymous fixed differences between the C57BL/6J and DBA/2J founder strains: *Ogg1*, a glycosylase that excises the oxidative DNA lesion 8-oxoguanine (8-oxoG) [4], and *Mbd4*, a glycosylase that can bind to methylated CpG sites and remove mispaired thymine nucleotides opposite spontaneously deaminated CpGs.

Missense mutations and loss-of-heterozygosity in *Ogg1* have been associated with increased risk of human cancer [28,29], and copy-number losses of either *Ogg1* or *Mutyh* are linked to elevated rates of spontaneous C>A mutation in human neuroblastoma [30]. Although *Ogg1* is a member of the same base-excision repair pathway as *Mutyh* (the protein-coding gene we previously implicated as harboring mutator alleles at the locus on chromosome 4), a number of lines of evidence suggest that the p.Asp129Asn missense mutation in *Mbd4* is the more compelling candidate mutator allele on chromosome 6. Unlike the *Ogg1* p.Thr95Ala mutation, p.Asp129Asn occurs at an amino acid residue within the *Mbd4* methyl-CpG binding domain that is well-conserved across mammalian species and predicted to be deleterious by *in silico* tools like SIFT [31] (Table 1). A missense mutation that affects the homologous amino acid in humans (p.Asp142Gly in GRCh38/hg38) is present on a single haplotype in the Genome Aggregation Database (gnomAD) [32] and is predicted by SIFT and Polyphen [33] to be "deleterious" and "probably_damaging" in human genomes, respectively. A recent study also identified a homozygous frameshift mutation in the primate homolog of *MBD4* that causes a hypermutator phenotype in the maternal germline [34], though this phenotype primarily comprises C>T mutations at CpG and CpA sites. Perhaps most intriguingly, loss-of-function mutations in *Mbd4* can sensitize cells to further DNA damage and prevent DNA damage-induced apoptosis [35,36].

We outline a plausible model of epistasis between *Mutyh* and *Mbd4* alleles in the *Supplementary Information*, but are unable to conclusively determine that either *Mbd4* or *Ogg1* harbors the causal variant underlying the observed C>A mutator phenotype. We anticipate that future experimental evidence will help pinpoint a causal allele and provide insight into the mechanism of epistasis between alleles at the loci on chromosome 4 and 6 in the BXDs.

Potential roles of structural variation and mobile elements as mutator alleles

Although we believe that *Mbd4* and *Ogg1* are the most likely candidate genes to explain the additional C>A mutator phenotype in the BXDs, we cannot conclusively determine that either the p.Asp129Asn or p.Thr95Ala missense mutation is a causal allele. We previously hypothesized that *Mutyh* missense mutations on *D* haplotypes were responsible for the large-effect C>A mutator phenotype we observed in the BXDs [2]. Using high-quality long-read assemblies of inbred laboratory strains, another group recently identified a ~5 kbp mobile element insertion (MEI) within the first intron of *Mutyh* [37] that is present on *D* haplotypes and absent from *B* haplotypes. This MEI is associated with significantly reduced expression of *Mutyh* in embryonic stem cells from laboratory strains, and may in fact underlie the previous C>A germline mutator phenotype in the BXDs. Although we did not find compelling evidence that structural variants were responsible for the novel C>A mutator phenotype observed in this study, it remains plausible that large SVs or MEIs contribute to mutation spectrum evolution in mammalian genomes.

Strengths and limitations of the inter-haplotype distance approach

The inter-haplotype distance (IHD) method has a number of advantages over traditional quantitative trait locus (QTL) mapping for detecting alleles that affect elements of the k -mer mutation spectrum. Using simulated data, we found that both IHD and QTL mapping had similar power to detect mutators that augment the rates of specific 1-mer and 3-mer mutation types (Figure 1-figure supplement 2). Nonetheless, only IHD was able to discover the mutator locus on chromosome 6 in the BXDs, demonstrating that it outperforms QTL mapping in certain experimental systems, and can reveal previously undiscovered signals of mutator alleles. One benefit of the IHD approach is that it obviates the need to perform separate association tests for every possible k -mer mutation type, and therefore the need to adjust significance thresholds for multiple tests. Since IHD compares the complete mutation spectrum between haplotypes that carry either allele at a site, it would also be well-powered to detect a mutator allele that exerted a coordinated effect on multiple k -mer mutation types (e.g., increased the rates of both C>T and C>A mutations).

However, the IHD method suffers a handful of drawbacks when compared to QTL mapping. Popular QTL mapping methods (such as R/qtld [38]) use linear models to test associations between genotypes and phenotypes, enabling the inclusion of additive and interactive covariates, as well as kinship matrices, in QTL scans. Although we have included simple methods to account for inter-sample relatedness in the IHD approach, they are not as robust or flexible as similar methods in QTL mapping software. Additionally, the IHD method assumes that mutator alleles affect a subset of k -mer mutation types, rather than all mutation types equally. If a mutator allele increased the rates of all mutation types equally on haplotypes that carried it, IHD would be unable to detect it.

Discovering mutator alleles in other systems

Mutator alleles likely contribute to variation in mutation rates and spectra across the tree of life. In two natural isolates of *Saccharomyces cerevisiae*, nonsynonymous variation in *OGG1* causes a substantial increase in the C>A *de novo* mutation rate [39]. Recent analyses have suggested that mutator alleles and/or environmental mutagens have shaped mutation rate evolution both in human

genomes [40] and more broadly during great ape evolution [41]. The heritability of paternal *de novo* mutation counts in the human germline has also been estimated to be between 10 and 20%, demonstrating a contribution of genetic factors to germline mutation rates [42]. However, mutator discovery remains challenging in mammalian genomes.

Recently, thousands of human pedigrees have been sequenced in an effort to precisely estimate the rate of human *de novo* germline mutation [6,7]. Selection on germline mutator alleles will likely prevent large-effect mutators from reaching high allele frequencies; as the number of active mutators increases, however, it becomes much more likely that a subset will be detectable by sequencing a sufficient number of human trios [43]. Current estimates of power to detect germline mutators in human pedigrees generally assume that mutators affect all mutation types equally, and that methods for mutator discovery will rely on identifying haplotypes with excess total mutation counts [43]. Our results in the BXDs suggest that germline mutators often exert their effects on a small number of k -mer mutation types, and may be far more amenable to detection by analyzing mutation spectra instead.

Materials and Methods

Identifying *de novo* germline mutations in the BXD RILs

The BXD resource currently comprises a total of 152 recombinant inbred lines (RILs). RILs were derived from either F2 or advanced intercrosses, and subsequently inbred by brother-sister mating for up to 180 generations [3]. BXDs were generated in distinct breeding “epochs,” which were each initiated with a distinct cross of C57BL/6J and DBA/2J parents; epochs 1, 2, 4, and 6 were derived from F2 crosses, while epochs 3 and 5 were derived from advanced intercrosses [3]. Previously, we analyzed whole-genome sequencing data from the BXDs and identified candidate *de novo* germline mutations in each line [2]. A detailed description of the methods used for DNA extraction, sequencing, alignment, and variant processing, as well as the characteristics of the *de novo* mutations, are available in a previous manuscript [2].

Briefly, we identified private single-nucleotide mutations in each BXD that were absent from all other RILs, as well as from the C57BL/6J and DBA/2J parents. We required each private variant to be meet the following criteria:

- genotyped as either homozygous or heterozygous for the alternate allele, with at least 90% of sequencing reads supporting the alternate allele
- supported by at least 10 sequencing reads
- Phred-scaled genotype quality of at least 20
- must not overlap regions of the genome annotated as segmental duplications or simple repeats in GRCm38/mm10
- must occur on a parental haplotype that was inherited by at least one other BXD at the same locus; these other BXDs must be homozygous for the reference allele at the variant site

A new approach to discover germline mutator alleles

Calculating inter-haplotype distance

Using the existing catalog of *de novo* germline mutations in the BXDs, we developed a new approach to discover loci that affect the germline *de novo* mutation spectrum in biparental RILs (Figure 1).

We assume that a collection of haplotypes has been genotyped at informative markers, and that *de novo* germline mutations have been identified on each haplotype.

At each informative marker, we divide haplotypes into two groups based on the parental allele that they inherited. We then compute a k -mer mutation spectrum using the aggregate mutation counts in each haplotype group. The k -mer mutation spectrum contains the frequency of every possible k -mer mutation type in a collection of mutations, and can be represented as a vector of size $6 \times 4^{k-1}$ after collapsing by strand complement. For example, the 1-mer mutation spectrum is a 6-element vector that contains the frequencies of C>T, C>G, C>A, A>G, A>T, and A>C mutations. Since C>T transitions at CpG nucleotides are often caused by a distinct mechanism (spontaneous deamination of methylated cytosine), we expand the 1-mer mutation spectrum to include a separate category for CpG>TpG mutations [44].

At each marker, we then compute the cosine distance between the two aggregate spectra. The cosine distance between two vectors \mathbf{A} and \mathbf{B} is defined as

$$D^C = 1 - \frac{\mathbf{A} \cdot \mathbf{B}}{\|\mathbf{A}\| \|\mathbf{B}\|}$$

where $\|\mathbf{A}\|$ and $\|\mathbf{B}\|$ are the L^2 (or Euclidean) norms of \mathbf{A} and \mathbf{B} , respectively. The cosine distance metric has a number of favorable properties for comparing mutation spectra. Since cosine distance does not take the magnitude of vectors into account, it can be used to compare two spectra with unequal total mutation counts (even if those total counts are relatively small). Additionally, by calculating the cosine distance between mutation spectra, we avoid the need to perform separate comparisons of mutation counts at each individual k -mer mutation type.

Inspired by methods from QTL mapping [38,45], we use permutation tests to establish genome-wide cosine distance thresholds. In each of N permutation trials, we randomly shuffle the per-haplotype mutation data such that haplotype labels no longer correspond to the correct mutation counts. Using the shuffled mutation data, we perform a genome-wide scan as described above, and record the maximum cosine distance observed at any locus. After N permutations (usually 10,000), we compute the $1 - p$ percentile of the distribution of maximum statistics, and use that percentile value as a genome-wide significance threshold (for example, at $p = 0.05$).

Accounting for relatedness between strains

We expect each BXD RIL to derive approximately 50% of its genome from C57BL/6J and 50% from DBA/2J. As a result, every pair of RILs will likely be identical-by-descent (IBD) at a fraction of genotyped markers. Pairs of more genetically similar BXDs may also have more similar mutation spectra, potentially due to shared polygenic effects on the mutation process. Therefore, at a given marker, if the BXD RILs that inherited D haplotypes are more genetically dissimilar from the RILs that inherited B haplotypes (considering all loci throughout the genome), we might expect the aggregate mutation spectra in the two groups to also be more dissimilar.

We implemented a simple approach to account for these potential issues of relatedness. At each marker g_i , we divide BXD haplotypes into two groups based on the parental allele they inherited. As before, we first compute the aggregate mutation spectrum in each group of haplotypes and calculate the cosine distance between the two aggregate spectra (D_i^C). Then, within each group of haplotypes, we calculate the allele frequency of the D allele at every marker along the genome to obtain a vector

of length n , where n is the number of genotyped markers. To quantify the genetic similarity between the two groups of haplotypes, we calculate the Pearson correlation coefficient r_i between the two vectors of marker-wide D allele frequencies.

Put another way, at every marker g_i along the genome, we divide BXD haplotypes into two groups and compute two metrics: D_i^C (the cosine distance between the two groups' aggregate spectra) and r_i (the correlation between genome-wide D allele frequencies in the two groups). To control for the potential effects of genetic similarity on cosine distances, we regress $(D_1^C, D_2^C, \dots, D_n^C)$ on (r_1, r_2, \dots, r_n) for all n markers using an ordinary least-squares model. We then use the residuals from the fitted model as the “adjusted” cosine distance values for each marker. If genome-wide genetic similarity between haplotypes perfectly predicts cosine distances at each marker, these residuals will all be 0 (or very close to 0). If genome-wide genetic similarity has no predictive power, the residuals will simply represent the difference between the observed cosine distance at a single marker and the marker-wide mean of cosine distances.

Implementation and source code

The inter-haplotype distance method was implemented in Python, and relies heavily on the following Python libraries: `numpy`, `pandas`, `matplotlib`, `scikit-learn`, `pandera`, `seaborn`, and `numba` [46,47,48,49,50,51,52].

Additional documentation is available on GitHub (<https://github.com/quinlan-lab/proj-mutator-mapping>), along with a reproducible Snakemake [53] workflow for running the method from start to finish using the BXDs (including downloading the mutation data, downloading genotypes, and running a genome-wide distance scan).

Simulations to assess the power of the inter-haplotype distance approach

We performed a series of simple simulations to estimate our power to detect alleles that affect the germline mutation spectrum in biparental RILs using the inter-haplotype distance method.

Simulating genotypes

First, we simulate genotypes on a population of haplotypes at a collection of sites. We define a matrix G of size (s, h) , where s is the number of sites and h is the number of haplotypes. We assume that every site is biallelic, and that the minor allele frequency at every site is 0.5. For every entry $G_{i,j}$, we take a single draw from a uniform distribution in the interval $[0.0, 1.0]$. If the value of that draw is less than or equal to 0.5, we assign the value of $G_{i,j}$ to be 1. Otherwise, we assign the value of $G_{i,j}$ to be 0.

Defining expected mutation type probabilities

Next, we define a vector of 1-mer mutation probabilities:

$$P = (0.29, 0.17, 0.12, 0.075, 0.1, 0.075, 0.17)$$

These probabilities sum to 1 and roughly correspond to the expected frequencies of C>T, CpG>TpG, C>A, C>G, A>T, A>C, and A>G *de novo* germline mutations in mice, respectively [10]. If we are simulating the 3-mer mutation spectrum, we modify the vector of mutation probabilities P to be length 96, and assign every 3-mer mutation type a value of $\frac{P_c}{16}$, where P_c is the probability of the

"central" mutation type associated with the 3-mer mutation type. In other words, each of the 16 possible NCN>NTN 3-mer mutation types would be assigned a mutation probability of $\frac{P_c}{16} = \frac{0.46}{16} = 0.02875$. We then generate a vector of lambda values by scaling the mutation probabilities by the number of mutations we wish to simulate (m):

$$\lambda = Pm$$

We also create a second vector of lambda values (λ'), in which we multiply the λ value of a single mutation type by the mutator effect size e .

In our simulations, we assume that genotypes at a single site (the "mutator locus") are associated with variation in the mutation spectrum. That is, at a single site s_i , all of the haplotypes with 1 alleles should have elevated rates of a particular mutation type and draw their mutation counts from λ' , while all of the haplotypes with 0 alleles should have "wild-type" rates of that mutation type and draw their mutation counts from λ . We therefore pick a random site s_i to be the "mutator locus," and identify the indices of haplotypes in G that were assigned 1 alleles at s_i . We call these indices h_{mut} .

Simulating mutation spectra

To simulate the mutation spectrum on our toy population of haplotypes, we define a matrix C of size (h, n) , where $n = 6 \times 4^{k-1}$ (or if $k = 1$ and we include CpG>TpG mutations, $6 \times 4^{k-1} + 1$).

Then, we populate the matrix C separately for *mutator* and *wild-type* haplotypes. For every row i in the matrix (i.e., for every haplotype), we first ask if i is in h_{mut} (that is, if the haplotype at index i was assigned a 1 allele at the "mutator locus"). If so, we set the values of C_i to be the results of a single Poisson draw from λ' . If row i is not in h_{mut} , we set the values of C_i to be the results of a single Poisson draw from λ .

Assessing power to detect a simulated mutator allele using IHD

For each combination of parameters (number of simulated haplotypes, number of simulated markers, mutator effect size, etc.), we run 100 independent trials. In each trial, we simulate the genotype matrix G and the mutation counts C . We calculate a "focal" cosine distance as the cosine distance between the aggregate mutation spectra of haplotypes with either genotype at s_i (the site at which we artificially simulated an association between genotype and mutation spectrum variation). We then perform an inter-haplotype distance scan using $N = 1,000$ permutations. If fewer than 5% of the N permutations produced a cosine distance greater than or equal to the focal distance, we say that the approach successfully identified the mutator allele in that trial.

Assessing power to detect a simulated mutator allele using quantitative trait locus (QTL) mapping

Using simulated data, we also assessed the power of traditional quantitative trait locus (QTL) mapping to detect a locus associated with mutation spectrum variation. As described above, we simulated both genotype and mutation spectra for a population of haplotypes under various conditions (number of mutations per haplotype, mutator effect size, etc.). Using those simulated data, we used R/qtL2 [38] to perform a genome scan for significant QTL as follows; we assume that the simulated genotype markers are evenly spaced (in physical Mbp coordinates) on a single chromosome. First, we calculate the fraction of each haplotype's *de novo* mutations that belong to each of the $6 \times 4^{k-1}$ possible k -mer mutation types. We then convert the simulated genotypes at each marker to genotype probabilities using the `calc_genoprob` function in R/qtL2, with `map_function = "c-f"` and

`error_prob = 0`. For every k -mer mutation type, we use genotype probabilities and per-haplotype mutation fractions to perform a scan for QTL with the `scan1` function; to make the results more comparable to those from the IHD method, we do not include any covariates in these QTL scans. We then use the `scan1perm` function to perform 1,000 permutations of the per-haplotype 1-mer mutation fractions and calculate log-odds (LOD) thresholds for significance. We consider the QTL scan to be “successful” if it produces a LOD score above the significance threshold (defined using $\alpha = \frac{0.05}{6 \times 4^{k-1}}$) for the marker at which we simulated an association with mutation spectrum variation.

Note: In our simulations, we augment the mutation rate of a single k -mer mutation type on haplotypes carrying the simulated mutator allele. However, in an experimental setting, we would not expect to have *a priori* knowledge of the mutation type affected by the mutator. Thus, by using an alpha threshold of 0.05, we would likely over-estimate the power of QTL mapping for detecting the mutator. Since we would need to perform 7 separate QTL scans (one for each 1-mer mutation type) in an experimental setting, we calculate QTL LOD thresholds at a Bonferroni-corrected alpha value of $\alpha = \frac{0.05}{6 \times 4^{k-1}}$.

Applying the inter-haplotype distance method to the BXDs

We downloaded previously-generated BXD *de novo* germline mutation data from the GitHub repository associated with our previous manuscript, which was also archived at Zenodo [2,54,55], and downloaded a CSV file of BXD genotypes at 7,320 informative markers from GeneNetwork [21,56]. We also downloaded relevant metadata about each BXD RIL from the manuscript describing the updated BXD resource [3]. These files are included in the GitHub repository associated with this manuscript.

As in our previous manuscript [2], we included mutation data from a subset of the 152 BXDs in our inter-haplotype distance scans. We removed any BXDs that had been inbred for fewer than 20 generations, as it takes approximately 20 generations of strict brother-sister mating for an RIL genome to become >98% homozygous [57]. As a result, any potential mutator allele would almost certainly be either fixed or lost after 20 generations. If fixed, the allele would remain linked to any excess mutations it causes for the duration of subsequent inbreeding, and its effects would be detectable using our methods. Additionally, a strain only meets the canonical definition of “inbred” if it has been subject to brother-sister mating for at least 20 generations [58]. We also removed the BXD68 RIL from our genome-wide scans, since we previously discovered a hyper-mutator phenotype in that strain; the C>A germline mutation rate in BXD68 is over 5 times the population mean, likely due to a private deleterious nonsynonymous mutation in *Mutyh* [2]. In total, we included 93 BXD RILs in our genome-wide scans.

We used Snakemake [59] to write a reproducible workflow for running the inter-haplotype distance method on the BXD dataset, which has been deposited in the GitHub repository associated with this manuscript.

Identifying candidate mutator alleles overlapping the chromosome 6 peak

We investigated the region implicated by our inter-haplotype distance approach on chromosome 6 by subsetting the joint-genotyped BXD VCF file (European Nucleotide Archive accession PRJEB45429 [60]) using `bcftools` [61]. We defined the candidate interval surrounding the cosine distance peak on chromosome 6 as +/- 5 Mbp from the genotype marker with the largest adjusted cosine distance value (`rs31001331`). To predict the functional impacts of both single-nucleotide variants and indels on splicing, protein structure, etc., we annotated variants in the BXD VCF using the following `snpEff` [62] command:

```
java -Xmx16g -jar /path/to/snpeff/jarfile GRCm38.75 /path/to/bxd/vcf > /path/to/uncompressed/output/vcf
```

and used `cyvcf2` [63] to iterate over the annotated VCF file in order to identify nonsynonymous fixed differences between the parental C57BL/6J and DBA/2J strains.

Extracting mutation signatures

We used SigProfilerExtractor (v.1.1.21) [25] to extract mutation signatures from the BXD mutation data. After converting the BXD mutation data to the “matrix” input format expected by SigProfilerExtractor, we ran the `sigProfilerExtractor` method as follows:

```
# install the mm10 mouse reference data
genInstall.install('mm10')

# run mutation signature extraction
sig.sigProfilerExtractor(
  'matrix',
  /path/to/output/directory,
  /path/to/input/mutations,
  maximum_signatures=5,
  nmf_replicates=50,
  opportunity_genome="mm10",
)
```

Comparing mutation spectra between Mouse Genomes Project strains

We downloaded mutation data from a previously published analysis [16] (Supplementary File 1, Excel Table S3) that identified strain-private mutations in 29 strains that were originally whole-genome sequenced as part of the Sanger Mouse Genomes (MGP) project [23]. When comparing counts of each mutation type between MGP strains that harbored either *D* or *B* alleles at the chromosome 4 or chromosome 6 mutator loci, we adjusted mutation counts by the number of callable A, T, C, or G nucleotides in each strain as described previously [2].

Querying GeneNetwork for eQTLs at the mutator locus

We used the online GeneNetwork resource [21], which contains array- and RNA-seq-derived expression measurements in a wide variety of tissues, to find *cis*-eQTLs for the DNA repair genes we implicated under the cosine distance peak on chromosome 6. On the GeneNetwork homepage (genenetwork.org), we selected the “BXD Family” **Group** and used the **Type** dropdown menu to select each of the specific expression datasets described in Table 2. In the **Get Any** text box, we then entered the listed gene name and clicked **Search**. After selecting the appropriate trait ID on the next page, we used the **Mapping Tools** dropdown to run Haley-Knott regression [64] with the following parameters: WGS-based marker genotypes, 1,000 permutations for LOD threshold calculations, and controlling for BXD genotypes at the `rs32497085` marker.

The exact names of the expression datasets we used for each tissue are shown in Table 2 below:

Table 2: Names of gene expression datasets used for each tissue type on GeneNetwork

Tissue name	Complete name of GeneNetwork expression data	GeneNetwork trait ID
Kidney	Mouse kidney M430v2 Sex Balanced (Aug06) RMA	1448815_at
Gastrointestinal	UTHSC Mouse BXD Gastrointestinal Affy MoGene 1.0 ST Gene Level (Apr14) RMA	10540639
Hematopoietic stem cells	UMCG Stem Cells ILM6v1.1 (Apr09) transformed	ILM1940279
Hematopoietic progenitor cells	UMCG Progenitor Cells ILM6v1.1 (Apr09) transformed	ILM1940279
Spleen	UTHSC Affy MoGene 1.0 ST Spleen (Dec10) RMA	10540639
Liver	UTHSC BXD Liver RNA-Seq Avg (Oct19) TPM Log2	ENSMUST00000032406
Heart	NHLBI BXD All Ages Heart RNA-Seq (Nov20) TMP Log2 **	ENSMUSG00000030271
Eye	UTHSC BXD All Ages Eye RNA-Seq (Nov20) TPM Log2 **	ENSMUSG00000030271

Calculating the frequencies of candidate mutator alleles in wild mice

To determine the frequencies of the *Ogg1* p.Thr95Ala and *Mbd4* p.Asp129Asn mutations in other populations of mice, we queried a VCF file containing genome-wide variation in 67 wild-derived mice from four species of *Mus* [26]. We calculated the allele frequency of each nonsynonymous mutation in each of the four species or subspecies (*Mus musculus domesticus*, *Mus musculus musculus*, *Mus musculus castaneus*, and *Mus spreitus*), including genotypes that met the following criteria:

- supported by at least 10 sequencing reads
- Phred-scaled genotype quality of at least 20

Testing for epistasis between the two mutator loci

To test for the presence of epistasis between the mutator loci on chromosome 4 and chromosome 6, we modeled C>A mutation rates in the BXDs as a function of genotypes at either locus. Specifically, we tested for statistical interaction between genotypes by fitting a generalized linear model in the R statistical language as follows:

```
m1 <- glm(Count ~ offset(log(ADJ_AGE)) + Genotype_A * Genotype_B, data =  
  data, family=poisson())
```

In this model, `Count` is the count of C>A *de novo* mutations observed in each BXD RIL. `ADJ_AGE` is the product of the number of “callable” cytosine/guanine nucleotides in each RIL (i.e., the total number of cytosines/guanines covered by at least 10 sequencing reads in the RIL) and the number of generations for which the RIL was inbred. We included the logarithm of `ADJ_AGE` as an “offset” in order to model the response variable as a rate rather than an absolute count; the BXDs differ in both

their durations of inbreeding and the proportions of their genomes that were sequenced to sufficient depth, which influences the number of mutations we observe in each RIL. The `Genotype_A` and `Genotype_B` terms represent the genotypes of BXDs at markers `rs52263933` and `rs31001331` (the markers with peak cosine distances on chromosomes 4 and 6 in the two inter-haplotype distance scans). Since each BXD is inbred for at least 20 generations, we considered genotypes at either locus to be binary ("B" or "D"). Using analysis of variance (ANOVA), we then compared the model including an interaction effect to a model including only additive effects:

```
m2 <- glm(Count ~ offset(log(ADJ AGE)) + Genotype_A + Genotype_B, data =  
          data, family=poisson())
```

```
anova(m1, m2, test="Chisq")
```

We tested for epistasis in the Sanger Mouse Genomes Project (MGP) strains using a nearly-identical approach. In this analysis, we fit two models as follows:

```
m1 <- glm(Count ~ offset(log(CALLABLE_C)) + Genotype_A * Genotype_B, data =  
          data, family=poisson())  
  
m2 <- glm(Count ~ offset(log(CALLABLE_C)) + Genotype_A + Genotype_B, data =  
          data, family=poisson())
```

where `Count` is the count of strain-private C>A mutations observed in each MGP strain [16]. The `CALLABLE_C` term represents the total number of cytosine and guanine nucleotides that were accessible for mutation calling in each strain, and the `Genotype_A` and `Genotype_B` terms represent MGP genotypes at `rs52263933` and `rs31001331`. We compared the two models using ANOVA as described above.

Since each BXD RIL derives approximately 50% of its genome from C57BL/6J and 50% from DBA/2J, we performed an additional test for epistasis that accounted for kinship between the BXD RILs. Using the `lme4` method from the `coxme` package [65] in the R statistical language, we fit a mixed effects model predicting C>A mutation fractions as a function of genotypes at both `rs52263933` and `rs31001331`, and included a pairwise kinship matrix as a random effect.

```
m = lme4(Fraction ~ Genotype_A * Genotype_B + (1| sample), data = data,  
          varlist = kinship_matrix)
```

The rows and columns of the kinship matrix were labeled with the `sample` name of each BXD, such that the `(1| sample)` term in the model captured the random effect of kinship. We calculated the `kinship_matrix` using the `calc_kinship` method from `R/qtl2` [38] as follows:

```

# read in the JSON-formatted file that directs R/qtl2 to sample
# genotypes, phenotypes, and covariates if applicable
bxd <- read_cross2("path/to/bxd.json")

# subset cross2 object to BXDs with C>A fractions in `data`
bxd <- bxd[data$sample, ]

# insert pseudomarkers into the genotype map
gmap <- insert_pseudomarkers(bxd$gmap, step = 0.2, stepwidth = "max")

# calculate QTL genotype probabilities
pr <- calc_genoprob(bxd, gmap, error_prob = 0.002, map_function = "c-f")

# calculate kinship between strains using all chromosomes
k <- calc_kinship(pr, "overall")

kinship_matrix = as.matrix(k)

```

Supplementary information

Chi-square statistics as an alternative to cosine distance in the inter-haplotype distance method

Although the current implementation of inter-haplotype distance (IHD) uses the cosine distance metric to compare aggregate mutation spectra at biallelic markers, we also explored the use of chi-square statistics to compare spectra.

As described in the *Materials and Methods*, we compute aggregate mutation spectra on haplotypes that inherited either parental allele at each biallelic marker. Rather than compute the cosine distance between those spectra, we can create a contingency table of size $(2, 6 \times 4^{k-1})$ (or size $(2, 6 \times 4^{k-1} + 1)$ if $k = 1$ and we're including CpG>TpG) that contains the aggregate k -mer mutation spectrum on haplotypes that inherited either parental allele. Using the contingency table, we then calculate a χ^2 test statistic; larger values of the χ^2 statistic suggest more “distance” between the aggregate mutation spectra on the two haplotypes.

We discovered the BXD mutator loci on chromosome 4 and 6 using either cosine distances or chi-square statistics to compare mutation spectra; however, using simulated data we found that the cosine distance metric was more sensitive for mutator allele detection across a broad range of parameter choices (Figure [1-figure supplement 3](#)).

Evidence for *Mbd4* as the causal gene underlying the chromosome 6 mutator locus

One of the key roles of *Mbd4* is to excise thymine nucleotides at G:T mispairs, which arise due to the spontaneous deamination of methylated CpGs. As a result, loss-of-function (LOF) mutations in *Mbd4* often lead to increased rates of CpG>TpG mutation [66]. Although LOF mutations in *Mbd4* are not known to cause C>A mutator phenotypes in mammalian cells, *Mbd4* is involved in a number of DNA

repair processes that may contribute to the mutator phenotype we observed in the BXDs. For example, bi-allelic LOF mutations in human *MBD4* underlie a neoplastic syndrome that closely mimics forms of familial adenomatous polyposis caused by LOF mutations in *MUTYH* [67]. Perhaps most intriguingly, LOF mutations in *Mbd4* can exacerbate the effects of exogenous DNA damage agents. Mouse embryonic fibroblasts that harbor homozygous LOF mutations in *Mbd4* fail to undergo apoptosis following treatment with a number of chemotherapeutics and mutagenic compounds [35]. Most of these exogenous mutagens cause DNA damage that is normally repaired by mismatch repair (MMR) machinery, but murine intestinal cells with biallelic LOF mutations in *Mbd4* also show a reduced apoptotic response to gamma irradiation, which is repaired independently of the MMR gene *Mlh1* [36]. Homozygous LOF mutations in *Mbd4* lead to accelerated intestinal tumor formation in mice that harbor an *Apc* allele that predisposes them to intestinal neoplasia [66], and mice with biallelic truncations of the *Mbd4* coding sequence exhibit modestly increased mutation rates in colon cancer cell lines, including increased C>A mutation rates in certain lines [68].

Mechanisms of epistasis between mutator alleles

Although both *Mutyh* and *Mbd4* are DNA glycosylases that participate in base-excision repair, they operate on distinct DNA lesions: *Mutyh* excises adenines paired with 8-oxoguanine, while *Mbd4* excises thymines opposite spontaneously deaminated CpGs. Given *Mbd4*'s role in suppressing DNA damage-induced apoptosis [35,36], we hypothesize that *D* alleles in *Mutyh* and *Mbd4* exhibit epistasis through the following mechanism in the BXD RILs.

In the absence of other defects in the DNA repair response, *D* alleles at *Mbd4* appear to have little or no detectable effect on *de novo* mutation rates (Figure 3). Since loss-of-function mutations in *Mbd4* often lead to increases CpG>TpG mutation rates [66], it is possible that the p.Asp129Asn missense mutation is not sufficient to cause such a phenotype in the BXDs. As we demonstrated in this and a previous manuscript, *D* alleles in *Mutyh* alone lead to significantly increased C>A mutation rates (Figure 3) [2]. In response to an accumulation of C>A mutations, a fraction of spermatagonial stem cells with *D* alleles at *Mutyh* may undergo apoptosis to prevent further unrepaired DNA damage. If those germline cells also harbor *D* alleles at *Mbd4*, they may not be able to arrest the cell cycle and complete apoptosis, allowing the effects of *D* alleles in *Mutyh* to exacerbate C>A mutation rates even further.

However, further experimental evidence will be required to identify the causal genes and mutations that underlie the observed signal of epistasis between mutator alleles.

Supplementary Figures

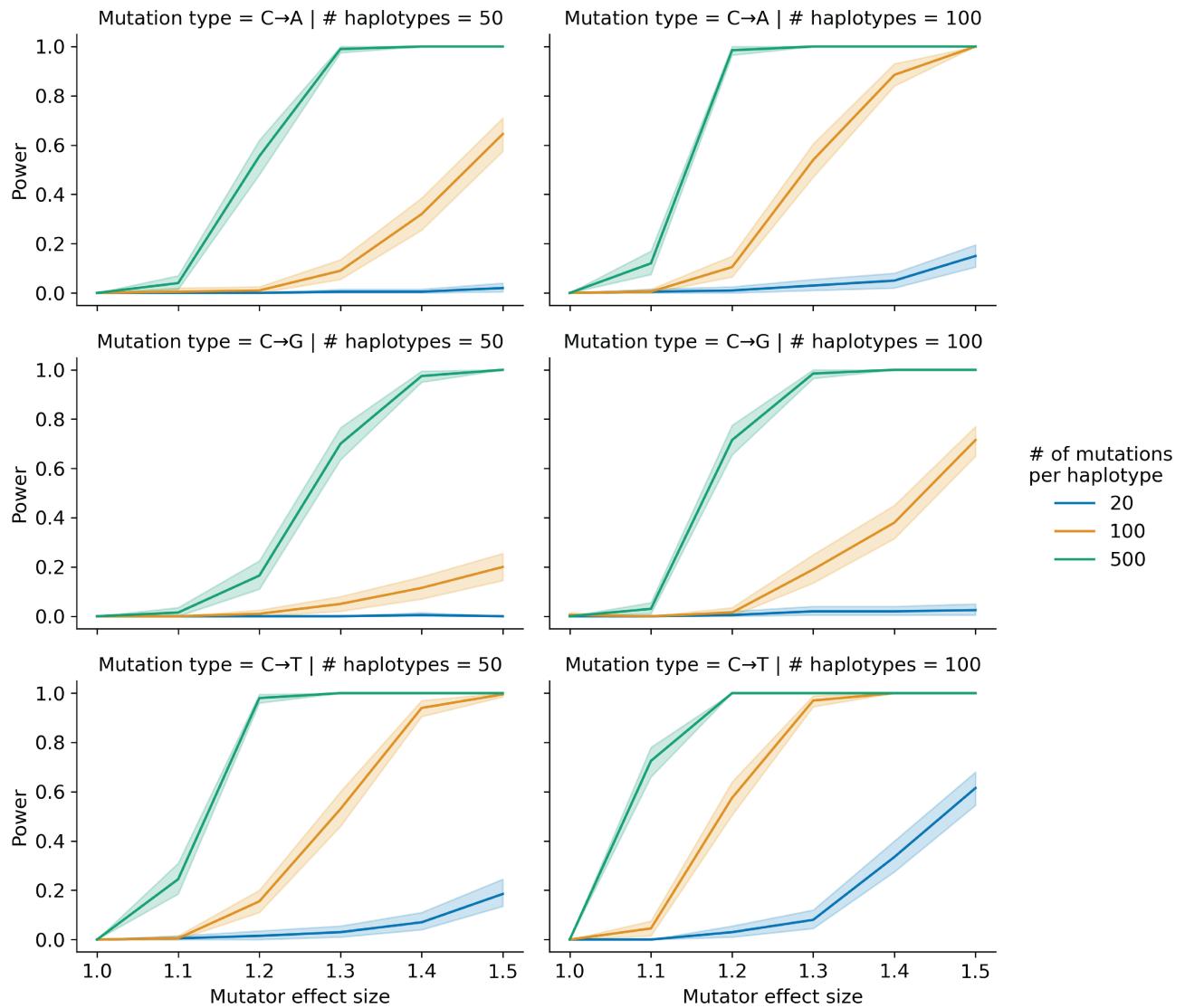


Figure 1-figure supplement 1: Simulations to assess the power of the inter-haplotype distance method. In each of 100 trials, we simulated genotypes at 1,000 biallelic loci on a toy population of either 50 or 100 haplotypes as follows. At every locus on every haplotype, we drew a single floating point value from a uniform distribution $[0, 1]$. If that value was less than or equal to 0.5, we set the allele to be "A"; otherwise, we set the allele to be "B". In each trial, we also simulated *de novo* germline mutations on the population of haplotypes, such that at a single locus g_i , we augmented the mutation rate of a particular k -mer by the specified effect size (an effect size of 1.5 indicates a 50% increase in the mutation rate) on haplotypes carrying "A" alleles. We then applied the inter-haplotype distance method to these simulated data and asked if the adjusted cosine distance at locus g_i was greater than expected by chance. Given a specific combination of parameters, the y-axis denotes the fraction of 100 trials in which the simulated mutator allele could be detected at a significance threshold of $p = 0.05$. Shaded areas indicate the standard deviation of that fraction across 100 simulations.

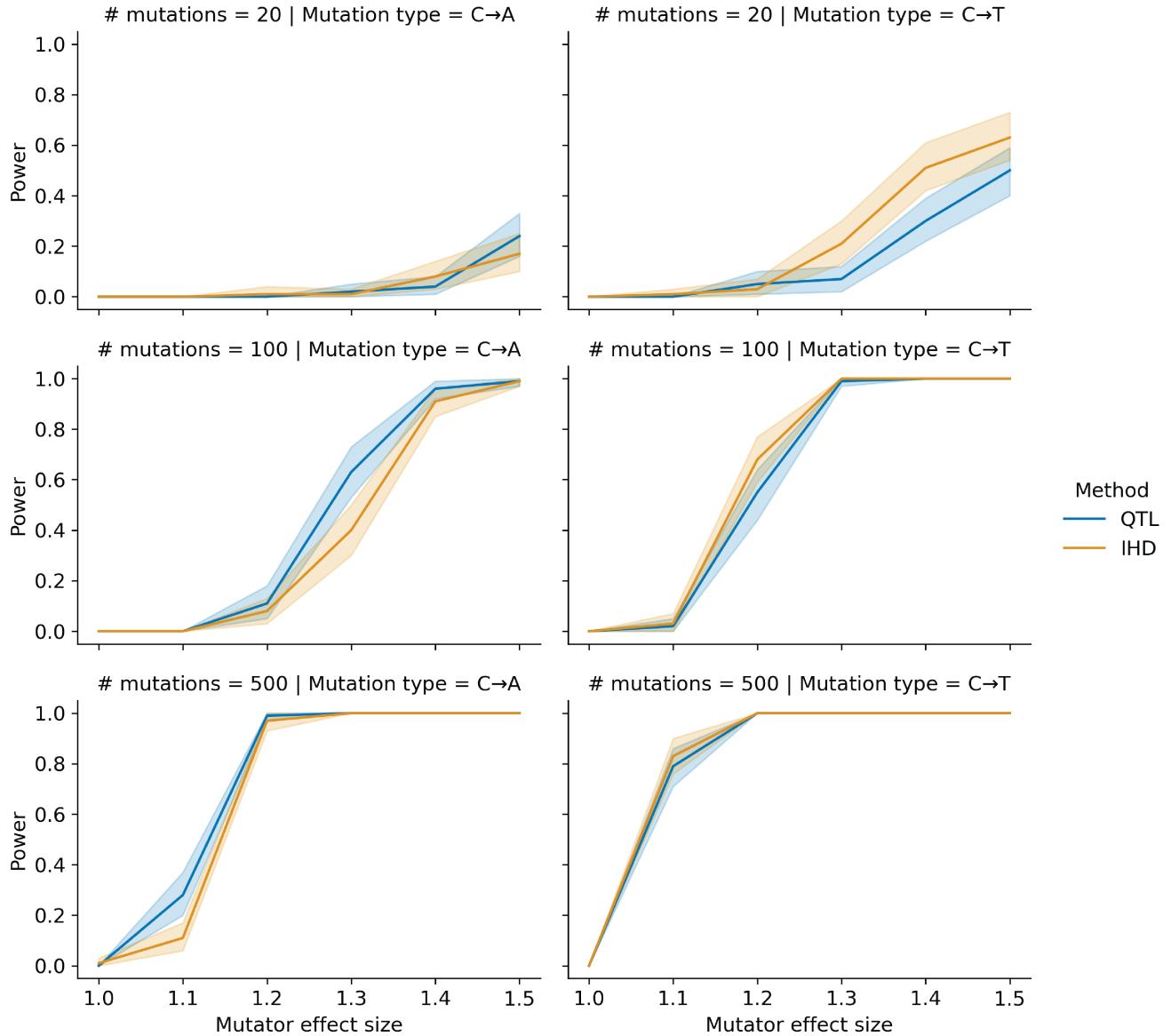


Figure 1-figure supplement 2: Comparing power between the inter-haplotype distance method and QTL mapping.

mapping. In each of 100 trials, we simulated genotypes at 1,000 biallelic loci on a toy population of 100 haplotypes as follows. At every locus on every haplotype, we drew a single floating point value from a uniform distribution $[0, 1]$. If that value was less than or equal to 0.5, we set the allele to be "A"; otherwise, we set the allele to be "B". In each trial, we also simulated *de novo* germline mutations on the population of haplotypes, such that at a single locus g_i , we augmented the rate of the specified mutation type by the specified effect size (an effect size of 1.5 indicates a 50% increase in the mutation rate) on haplotypes carrying "A" alleles. We then applied the inter-haplotype distance method to these simulated data and asked if the adjusted cosine distance at locus g_i was greater than expected by chance. Similarly, in each trial, we used R/qt12 to perform a genome scan for QTL and asked if the log-odds score at g_i was greater than expected by chance. Given a specific combination of parameters, the y-axis denotes the fraction of 100 trials in which the simulated mutator allele could be detected at a significance threshold of $p = 0.05$ (for IHD) or at an alpha of $\frac{0.05}{7}$ (for QTL mapping). Shaded areas indicate the standard deviation of that fraction across 100 simulations.

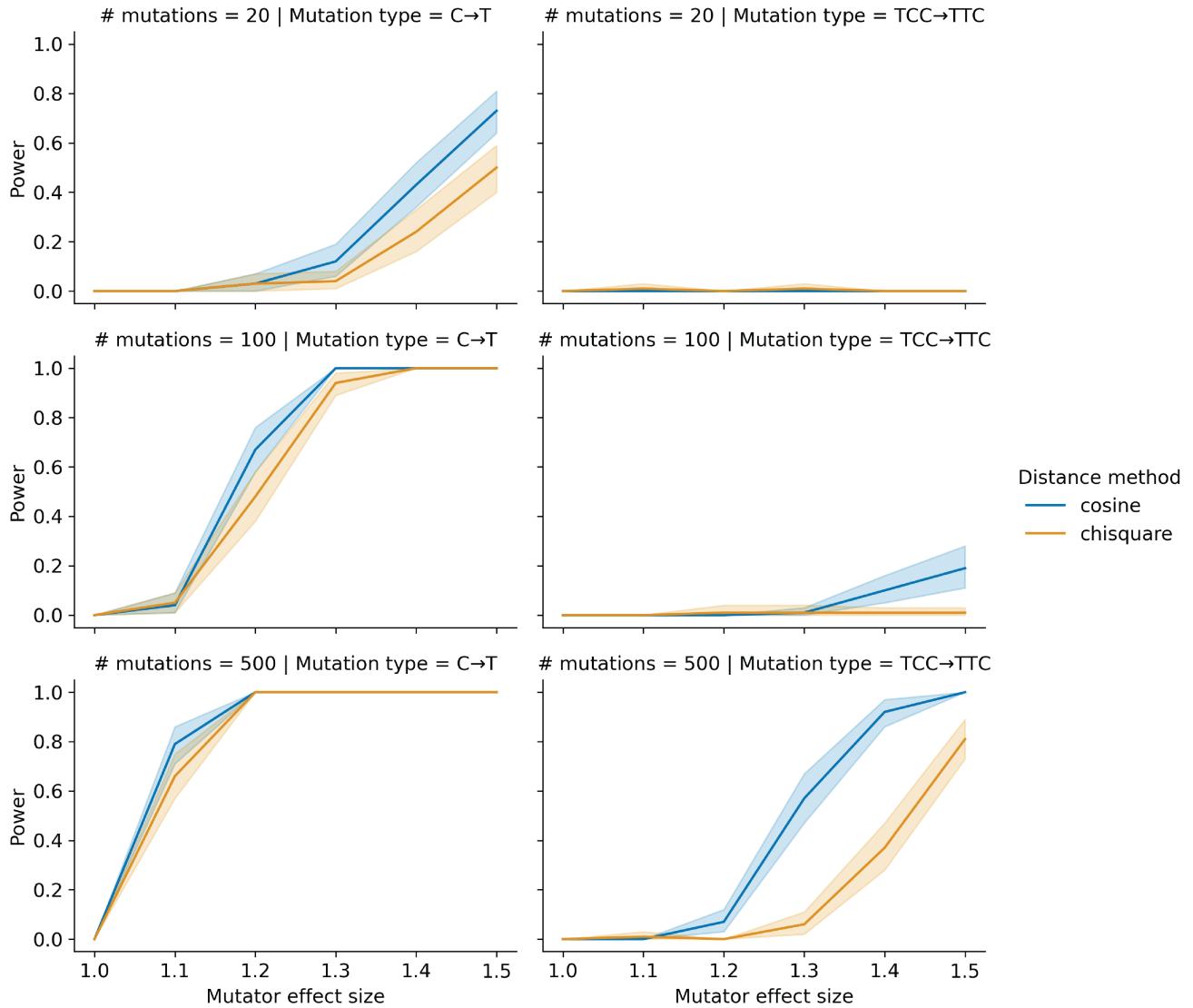


Figure 1-figure supplement 3: Comparing the power of cosine distance and chi-square statistics in the inter-haplotype distance method. In each of 100 trials, we simulated genotypes at 1,000 biallelic loci on a toy population of 100 haplotypes as follows. At every locus on every haplotype, we drew a single floating point value from a uniform distribution $[0, 1]$. If that value was less than or equal to 0.5, we set the allele to be "A"; otherwise, we set the allele to be "B". In each trial, we also simulated *de novo* germline mutations on the population of haplotypes, such that at a single locus g_i , we augmented the rate of the specified mutation type by the specified effect size (an effect size of 1.5 indicates a 50% increase in the mutation rate) on haplotypes carrying "A" alleles. We then applied the inter-haplotype distance method, using either cosine distance or chi-square statistics to compare aggregate mutation spectra at each marker. In each trial, we asked if the adjusted cosine distance or chi-square statistic at locus g_i was greater than expected by chance. Given a specific combination of parameters, the y-axis denotes the fraction of 100 trials in which the simulated mutator allele could be detected at a significance threshold of $p = 0.05$. Shaded areas indicate the standard deviation of that fraction across 100 simulations.

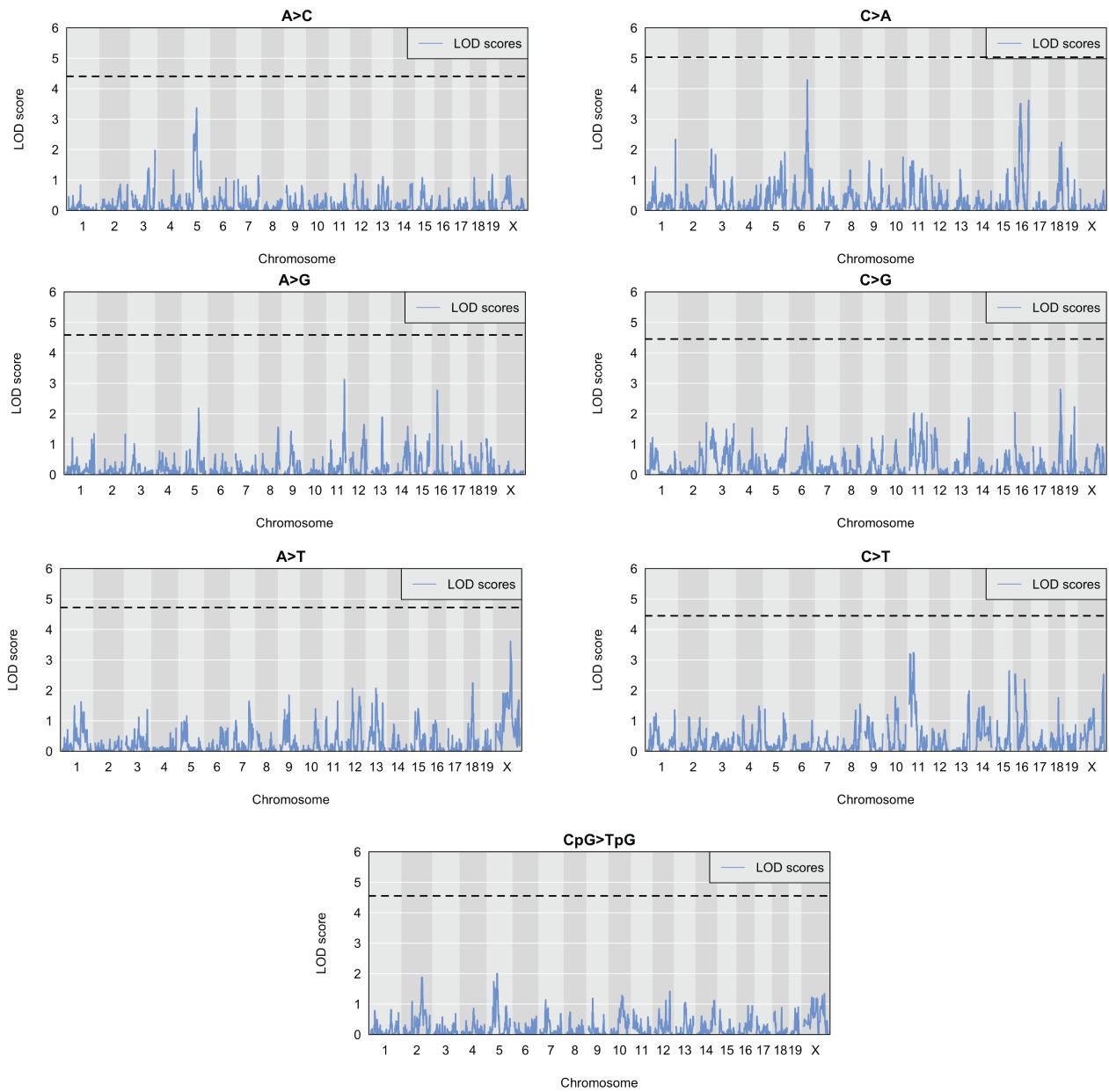


Figure 2-figure supplement 1: Quantitative trait locus scans for mutation spectrum phenotypes. Using the BXDs with D genotypes at rs52263933 (the marker with the highest cosine distance on chromosome 4), we used R/qt12 to perform QTL scans for the fraction of each 1-mer mutation type. Plots show the log-odds (LOD) score at every genotyped marker in blue; the dotted black line represents the genome-wide LOD significance threshold (established using 1,000 permutations at an alpha of $\frac{0.05}{7}$ to account for the fact that 7 separate association tests were performed.)

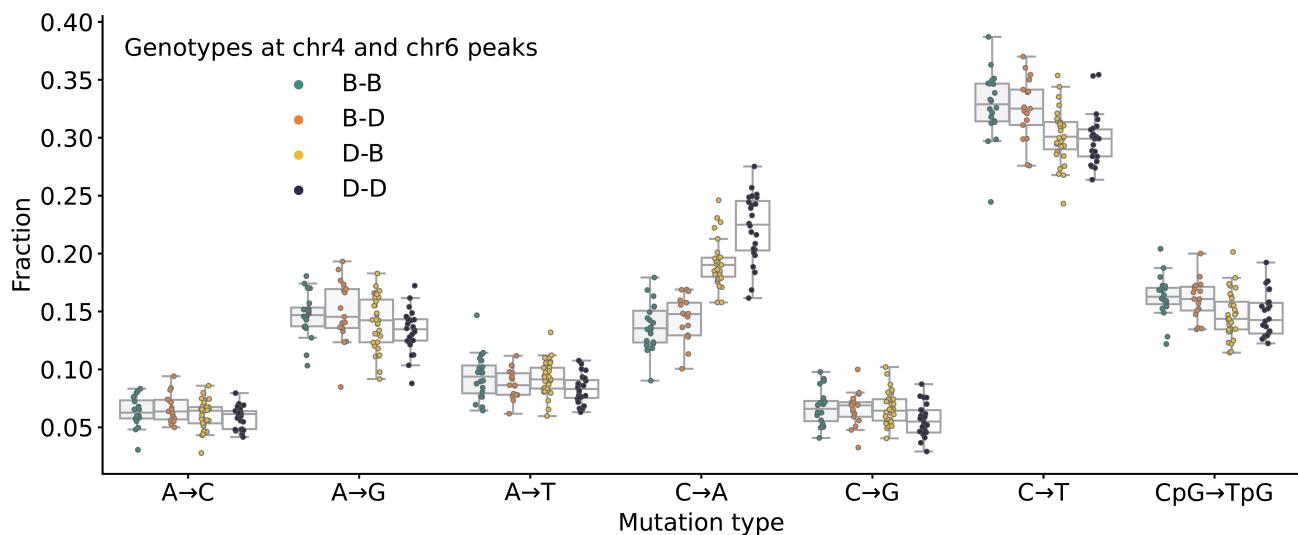


Figure 2—figure supplement 2: Mutation spectra comparison in BXD strains. Fractions of *de novo* germline mutations in BXDs with either D or B genotypes at markers rs52263933 and rs31001331, stratified by mutation type.

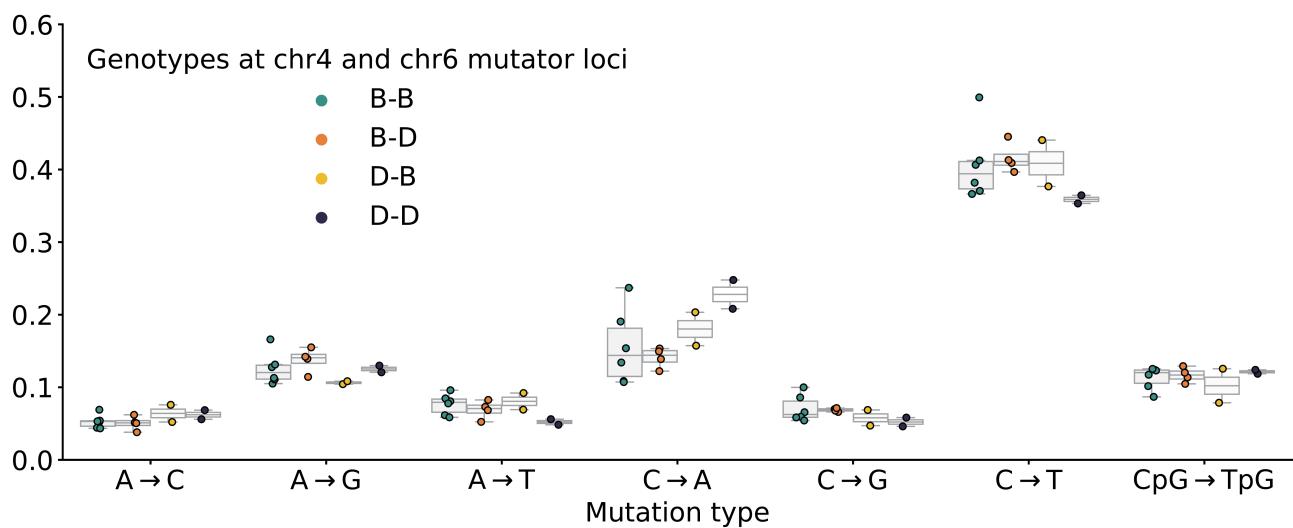


Figure 3—figure supplement 1: Mutation spectra comparison in Sanger Mouse Genomes Project strains. Fractions of *de novo* germline mutations in Sanger MGP strains with either D or B genotypes at markers rs52263933 and rs31001331, stratified by mutation type.

Supplementary Tables

Table supplement 1: Presence or absence of cis-eQTLs for *Ogg1* and *Mbd4* in various tissues identified using GeneNetwork.

Gene name	Tissue name	# BXDs with expression data	Top significant marker	LRS at top significant marker	Significant LRS threshold
<i>Ogg1</i>	Kidney	53	rsm100000041 88	52.25	17.82
<i>Ogg1</i>	Gastrointestinal	46	rsm100000034 41	23.39	16.09
<i>Ogg1</i>	Hematopoetic stem cells	22	-	-	16.45

Gene name	Tissue name	# BXDs with expression data	Top significant marker	LRS at top significant marker	Significant LRS threshold
<i>Ogg1</i>	Hematopoetic progenitor cells	23	-	-	18.52
<i>Ogg1</i>	Spleen	79	rsm100000034 18	-	17.51
<i>Ogg1</i>	Liver	50	rsm100000041 88	53.54	18.77
<i>Ogg1</i>	Heart	73	-	-	16.22
<i>Ogg1</i>	Eye	87	rsm100000041 94	23.05	16.96
<i>Mbd4</i>	Kidney	53	-	-	18.48
<i>Mbd4</i>	Gastrointestinal	46	-	-	15.97
<i>Mbd4</i>	Hematopoetic stem cells	22	-	-	18.12
<i>Mbd4</i>	Hematopoetic progenitor cells	23	-	-	18.55
<i>Mbd4</i>	Spleen	79	rsm100000041 99	21.42	16.99
<i>Mbd4</i>	Liver	50	-	-	16.15
<i>Mbd4</i>	Heart	73	-	-	17.17
<i>Mbd4</i>	Eye	87	-	-	16.63

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