

## Methods

# The Development of a Highly Informative Mouse Simple Sequence Length Polymorphism (SSLP) Marker Set and Construction of a Mouse Family Tree Using Parsimony Analysis

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To identify highly informative markers for a large number of commonly employed murine crosses, we selected a subset of the extant mouse simple sequence length polymorphism (SSLP) marker set for further development. Primer pairs for 314 SSLP markers were designed and typed against 54 inbred mouse strains. We designed new PCR primer sequences for the markers selected for multiplexing using the fluorescent dyes FAM, VIC, NED, and ROX. The number of informative markers for C57BL/6J × DBA/2J is 217, with an average spacing of 6.8 centiMorgans (cM). For all other pairs of strains, the mean number of informative markers per cross is 197.0 (SD 37.8) with a mean distance between markers of 6.8 cM (SD 1.1). To confirm map positions of the 224 markers in our set that are polymorphic between *Mus musculus* and *Mus spretus*, we used The Jackson Laboratory (TJL) interspecific backcross mapping panel (TJL BSS); 168 (75%) of these markers had not been previously mapped in this cross by other investigators, adding new information to this community map resource. With this large data set, we sought to reconstruct a phylogenetic history of the laboratory mouse using Wagner parsimony analysis. Our results are largely congruent with the known history of inbred mouse strains.

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Other investigators have developed simple sequence length polymorphism (SSLP) markers for use as genetic markers in many organisms. Short tandem repeat (STR), short tandem repeat polymorphism (STRP), and microsatellite are other names commonly used for this type of marker. A comprehensive genetic linkage map of the mouse has been constructed (Dietrich et al. 1996) using over 6000 SSLP markers with an average spacing of less than 1 cM. However, it is still difficult to identify sufficient markers that are informative for a specific pair of inbred strains, particularly if they are both derived from *Mus musculus domesticus*. Allele size data are available for only a limited number of inbred mouse strains (Schalkwyk et

al. 1999), which makes it difficult to construct a suitable panel of markers that are detectably polymorphic for a selected cross. In addition, many primer pairs have not been extensively characterized or do not yield robust PCR products.

Starting with the CA repeat-containing sequences identified by Dietrich et al. (1996), we assembled a set of marker loci that span the mouse genome and are informative in many common inbred strain combinations, and we designed new optimized primer sequences. Our primers were specifically designed to allow multiplexing by fluorescent dye color and product size. Our current level of multiplexing is 8–9 loci per capillary on the Applied Biosystems 3700 platform, producing ~750 genotypes per run. The forward primer of each pair was labeled on the 5' end with the fluorescent dyes FAM, VIC, or NED. We avoided CA sequences on the 3' end of the forward primers to reduce stuttering artifacts. All reverse primers contain the sequence GTGTCTT (5'–3') at their 5' ends to promote nontemplate-directed nucleotide addition

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(+A; Smith et al. 1995; Brownstein et al. 1996). This simplifies allele calling by reducing stuttering artifacts that arise during amplification of repeat-containing sequences. Theoretically, we could multiplex more loci per lane depending on the sensitivity of the detection platform and the robustness of the PCR reactions. We presently multiplex post-PCR by pooling different ratios of each PCR product.

In order to confirm the map positions identified by our newly designed PCR assays for this set of SSLP markers, we typed 94 animals from TjL BSS interspecific mapping panel [(C57BL/6J)Ei × SPRET/Ei]F1 × SPRET/Ei to place our markers in this existing comprehensive map that contained over 4500 mapped loci at the time of our analysis (Rowe et al. 1994).

SSLP markers are widely dispersed, selectively neutral, and their alleles typically differ by discrete steps (Weber and Wong 1993). For these reasons, they have proven useful for creating mouse family trees by parsimony analysis. To infer the degree of relatedness between inbred mouse strains, we used the allele size data for all 314 markers that we collected for 54 strains to construct a mouse family tree as described by Acthley and Fitch (1993) and Schalkwyk et al. (1999). We performed Wagner parsimony analysis using the MIX program of PHYLIP v. 3.57 (Felsenstein 1989).

## RESULTS AND DISCUSSION

### Mouse Linkage Mapping Set

We chose a target density of 10 cM for our marker set in order to conduct genome-wide screens for linkage analysis. We first selected ~160 markers from the Dietrich et al. (1996) set to maximize polymorphism for the cross C57BL/6J × DBA/2J. We then selected an additional ~160 markers to increase the map density and maximize polymorphism for other pairs of strains. We designed and tested a total of 323 primer pairs for 314 SSLP markers. Twenty-two primer pairs failed or performed poorly. Thirteen of these primer pairs were successfully redesigned, resulting in 305 usable markers. The remaining nine markers were replaced to fill any remaining gaps between markers that were greater than 10 cM. The total number of markers for which primers were successfully designed is 314. Map positions are based on stored values in the Mouse Genome Database (MGD; TjL) which are updated daily.

The result was a set of 314 SSLP markers, which we used to genotype a total of 54 inbred mouse strains. Graphical chromosome maps displaying the cM position of each marker are shown in Figure 1. The number of informative markers for the C57BL/6J × DBA/2J cross is 217, with an average spacing of 6.8 cM. This represents ~70% of the markers in the mapping set. The largest gap for this cross is 33 cM at the end of chromosome 5. For all other pairs of strains, the mean number of informative markers per cross is 197.0 (SD 37.8). The mean distance between markers is 6.8 cM (SD 1.1), and the mean number of gaps greater than 20 cM is 7.

A histogram showing the number of informative markers for each pair of mouse strains we evaluated is shown in Figure 2. The tail on the left side of the histogram displays crosses between closely related substrains. This includes a cluster of six 129 substrains. The genetic variability among 129 substrains was documented by Simpson et al. (1997), who observed differences as high as 22% for alleles at 86 SSLP loci in fifteen 129 substrains. We genotyped ~3.5-fold more markers

and observed differences of 16%–26% among six 129 substrains. We concur with Simpson et al. (1997) that the majority of this variability is most likely the result of unrecognized outcrossing rather than new mutations. The contamination of 129/SvJ has been reported elsewhere (Threadgill et al. 1997). Table 1 indicates the number of informative markers for crosses among 12 inbred mouse strains.

Current, detailed information about this marker set is publicly available on the Web at <http://www.cidr.jhmi.edu/mouse/mouse.html>. Investigators can submit a query for any pairwise combination of inbred strains to retrieve the informative markers for a cross, or download a spreadsheet with the locus names, map positions, and allele sizes for each marker for all 54 strains. We have incorporated other features into our Web site (<http://www.cidr.jhmi.edu/mouse/queries.html>), including a panel generator to group primer pairs by dye color and allele sizes for multiplex loading of gels or capillaries, and a query to identify markers for constructing speed congenics by marker-assisted selection (Wakeland et al. 1997).

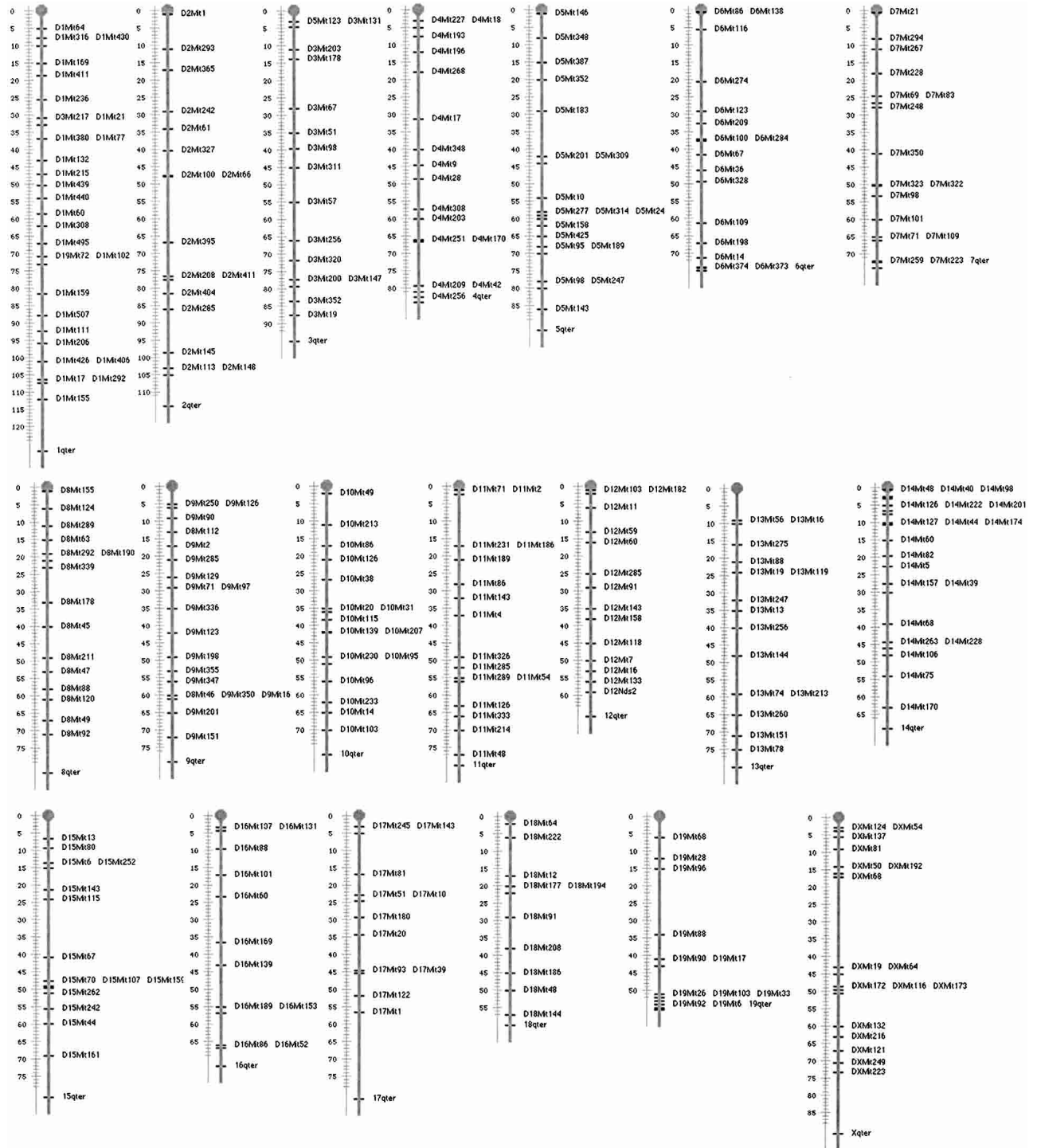
The panel generator allows the user to (1) select the maximum number of primer pairs in a panel, (2) select the maximum number of primer pairs with the same dye color in a panel, (3) increase the size ranges of each primer pair to avoid overlaps with other primer pairs, and (4) select the minimum number of base pairs between the size ranges of primer pairs with the same dye color in a panel. A list of all of the informative markers is displayed, along with map information, the spacing between markers, PCR performance, and comments about the primer pairs. The user can choose the best markers at the desired spacing to generate the panels. Panels are displayed on a new page with the dye color, size range, allele sizes, and recommended PCR conditions for each primer pair. The user can then click on a link that will display the part numbers for the primer pairs, which can be ordered directly from Applied Biosystems. These marker panels are specific for the amplicon sizes and dye colors in the primer pairs available from Applied Biosystems.

The speed congenics query generates a list of informative markers between the donor and recipient strains such that recipient allele sizes must be different from donor allele sizes. Investigators can select up to three donor strains and up to three recipient strains. We also included a query that only displays markers in which the allele sizes of all of the selected strains are different.

The current marker density is sufficient to conduct 20-cM genome-wide screens in mouse for linkage mapping studies (Silver 1995). For such scans we routinely type approximately 85 markers. We have used this marker set to type 1843 animals for five different studies involving seven different strains. Four studies were genome-wide screens for quantitative trait loci (QTLs), and the fifth study was the first phase of a marker-assisted breeding experiment. The cumulative average missing data rate we have achieved is 5.2%, and the cumulative average error rate as calculated from 5% blind duplicate samples run concurrently with each study is 0.1%.

The primer pairs we designed are different from the primer pairs designed by The Whitehead Institute/MIT Center for Genome Research (WICGR). Few, if any, of the allele sizes observed using these primers will be the same as those observed using the WICGR primers. However, the markers should remain informative for a pair of inbred mouse strains using either set of primers. We developed and characterized all of the markers using Applied Biosystems 377XL sequenc-

## Mouse SSLP and Family Tree by Parsimony Analysis

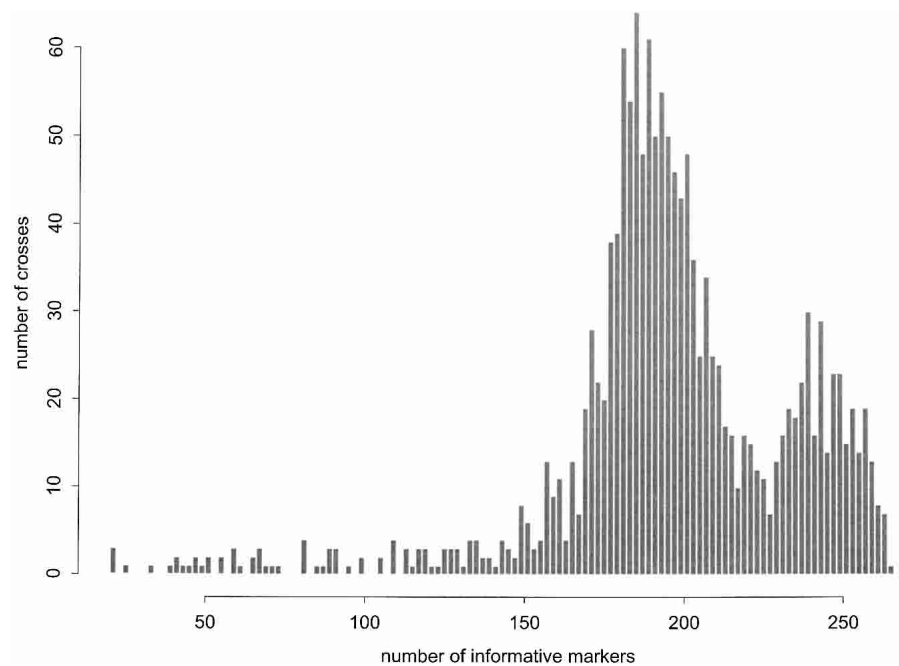


**Figure 1** Graphical chromosome maps displaying the map positions of 314 newly designed SSLP markers for fluorescent genotyping by PCR analysis. Map positions in centimorgans (cM) were obtained from the Mouse Genome Database (TJL) with the exception of *D3Mit217*, *D19Mit72*, *D8Mit46*, and *D8Mit112*, which are displayed relative to their map order on The Jackson Laboratory BSS Mapping Panel.

ers. Running the products on a different instrument platform or with a different size standard will often result in different observed allele sizes, but the size difference between strains should be constant.

### Backcross Mapping

We used TJL BSS interspecific backcross mapping panel to confirm the map positions of the markers in our set that were informative between *M. musculus* and *M. spretus*. Although 251



**Figure 2** Number of informative markers per cross. Pairwise combinations of 54 inbred mouse strains were analyzed to determine the number of markers that are polymorphic for each cross. The number of crosses are plotted against the number of informative markers for each pair of strains. The mean number of informative markers per cross for all pairs of strains analyzed is 197.0 (SD 37.8).

markers from our set are polymorphic for this strain combination, 27 of these could not be mapped due to allele dominance (primers designed to C57BL/6J sequence preferentially amplify only from the C57BL/6J allele template in the presence of a highly diverged *M. spretus* allele sequence). We also observed that a small percentage of marker loci fail to amplify for strains not belonging to the *M. m. domesticus* subgroup (data not shown).

We mapped the remaining 224 informative markers onto The Jackson Laboratory BSS Mapping Panel; 220 (98%) marker loci mapped to the expected chromosome location. One hundred and sixty-eight of these markers (75%) represent new loci that had not previously been mapped by other investigators onto the BSS panel, contributing new information to this comprehensive genetic map of the mouse. Detailed information for all loci mapped in this cross including

the mapping data reported here can be found at <http://www.jax.org/resources/documents/cmldata/bkmap/BSS.html>.

Four (2%) of our mapped marker loci did, however, map to unexpected positions. D3Mit217 and D19Mit72 mapped to chromosome 1, whereas D8Mit46 and D8Mit112 both mapped to chromosome 9. All of these new chromosomal assignments are corroborated by independent data sets from other laboratories submitted to The Jackson Laboratory Mapping Panels, using the interspecific backcross panels or the mouse T31 Radiation Hybrid mapping panel (<http://www.jax.org/resources/documents/cmldata/rhmap/RHIntro.html>) or both. Based on this evidence, these four markers have been renamed by the Mouse Locus Nomenclature Committee as of May 2002 as *D1Mit1000* for the old *D3Mit217*, *D1Mit1001* for the former *D19Mit72*, *D9Mit1000* for the locus previously named *D8Mit46*, and *D9Mit1001* for the old *D8Mit112*.

We compared our BSS mapping results to the composite maps available from the MGD (<http://www.informatics.jax.org>). We used the MGD values to generate the chromosome maps in Figure 1. Composite chromosome maps are longer than the BSS maps, because composite maps are based on many types of crosses and other mapping methods. Recombination frequencies vary from one cross to another, and the number of commonly mapped anchor markers in some combinations of crosses can be scarce, and because creating composite maps is an inherently inaccurate process, particularly for closely linked loci mapped in independent crosses or systems. In addition, any data errors expand the apparent distances between markers. Although these problems can and do lead to order inaccuracies in composite maps, we observe no differences between the order of our markers in the TJL BSS map and MGD composite maps.

**Table 1.** Number of Polymorphic Markers Between 12 Pairs of Inbred Mouse Strains

	DBA/2J	129P3/J	A/J	BALB/cJ	FVB/NJ	NOD/LtJ	CAST/Ei	C3H/HeJ	AKR/J	NZB/B1NJ	SPRET/Ei
DBA/2J											
129P3/J	181										
A/J	177	177									
BALB/cJ	172	186	109								
FVB/NJ	184	196	156	158							
NOD/LtJ	185	191	166	178	161						
CAST/Ei	259	251	263	262	260	255					
C3H/HeJ	163	193	134	158	171	185	259				
AKR/J	188	197	160	167	182	178	258	177			
NZB/B1NJ	200	194	210	196	192	193	263	220	200		
SPRET/Ei	234	231	236	240	238	236	238	234	241	243	
C57BL/6J	217	197	223	214	217	213	269	246	219	204	244



gains or losses of a single repeat unit (Weber and Wong 1993). However, multiple repeat units can be added or deleted between alleles, and sometimes the entire repeat is duplicated. Without a defined model for the process of addition or deletion of multiple repeat units, it is not clear how this process might affect the analysis. Larger data sets should minimize any distortion of results due to this type of variation. We coded allele sizes as  $n-1$  integers, where  $n$  is the number of alleles, and allele sizes differ by one step (i.e., two base pairs). This gives order to the mutational changes between alleles but does not assume that a particular allele is ancestral. We performed a Wagner parsimony analysis using the MIX program of PHYLIP v. 3.57 (Felsenstein 1989). User options were selected to generate an unrooted tree after the input order of allele sizes was randomized three times. No ancestral state for allele sizes was assumed for analysis. A consensus family tree is shown in Figure 3.

This tree has six major groups of branches, labeled 1 to 6. If recent inbreeding occurred between the strains, we would observe a net rather than a tree structure. Group 1 is comprised of Bagg albino derivatives and includes I/LnJ mice. Strong originally derived these mice from an unpedigreed stock. Group 2 is made up of strains derived from species of wild mice not belonging to *M. m. domesticus* (Bonhomme and Guenet 1996). Group 3 contains C57 and C58 animals from Miss Abbie Lathrop's stock, which unlike most "old" inbred strains carry a Y chromosome of Asian *M. m. musculus* origin (Tucker et al. 1992). Group 4 contains Castle's 129 and LP mice, which are known to be similar. 101H has a common origin with 129. Groups 3 and 4 appear to be parts of a large limb, and their members may be related. In light of the fact that both sets of animals were generated from coat color stocks provided by English fanciers, this seems reasonable (Potter 1978). Groups 5 and 6 hold animals that appear to derive from European ("Swiss") *M. m. domesticus*. BDP, P, and SM mice (Group 5) are grouped separately from DBA (Group 1). BDP was produced by crossing Strong's pink-eyed and Little's DBA. P was derived by Snell from a cross involving BDP. SM was produced by crossing seven strains, including DBA, and selecting for small body size. These strains appear to have diverged significantly from the Bagg albino group. The BUB/BnJ inbred strain (Group 6) originated from randomly bred albino mice of unknown ancestry.

We considered the possibility that our results would be biased by the selection of markers informative for the C57BL/6J  $\times$  DBA/2J strain combination. However, parsimony analysis considers specific mutation steps rather than the simple proportion of alleles that are the same size or different for a given marker. Further, it assumes that species and characters evolve independently. It does not assume that the ancestral state is known or that mutation in one direction is more probable than the other. Nonetheless, our results are largely congruent with the known history of inbred mouse strains (Festing 1996; Beck et al. 2000). Discretion should be used, however, when considering this information to ascertain the effects of genetic background when assessing phenotypes.

## METHODS

### Primers

Oligonucleotides were synthesized by Applied Biosystems. Primers were designed using DNA sequences from The Whitehead Institute/MIT Center for Genome Research (WICGR) Genetic Map of the Mouse Database Release 10 ([http://www.ensembl.org/Mus\\_musculus](http://www.ensembl.org/Mus_musculus)).

The forward primer of each pair was labeled on the 5' end with the fluorescent dye FAM, VIC, or NED. To reduce stuttering artifacts, CA dinucleotide sequences were avoided on the 3' end of the forward primers. All reverse primers contain the sequence GTGTCTT (5'-3') at their 5' ends to promote nontemplate-directed nucleotide addition (+A) to the PCR products. Primer sequences are proprietary. Primer pairs can be purchased individually or as a complete set directly from Applied Biosystems.

### DNA

Genomic DNA for all inbred mouse strains was obtained from The Jackson Laboratory with the exception of 129S2/SvPas (Charles River Laboratories) and 129S6/SvEv (Taconic). PL/J, LT/SvEv, C3H/HeN, C3H/HeSnJ, BALB/cByJ, and 129/Sv (agouti) DNA were a gift from E. Eicher and T. Golovkina (TJL). LG/J DNA was a gift from J. Cheverud and S. Cropp (Washington University). 101H DNA was a gift from P. Denny and A. Southwell (Harwell, UK). DNA from the BSS interspecific backcross panel [(C57BL/6)Ei  $\times$  SPRET/Ei]F1  $\times$  SPRET/Ei] was obtained from L. Rowe (TJL).

### PCR

PCR reactions were performed using PE 9700 thermal cyclers in a volume of 10  $\mu$ L. Final reaction conditions were 50 ng DNA, 250  $\mu$ M each dNTP, 2.5 mM MgCl<sub>2</sub>, 0.4  $\mu$ M each primer, and 0.25 U Taq Gold DNA Polymerase (Applied Biosystems). PCR was initiated with an automatic hot start for 15 min at 95°C to activate the Taq Gold polymerase. Standard cycling conditions were carried out at 94°C for 45 sec, 55°C for 60 sec, and 72°C for 60 sec (30 cycles) with a final extension at 72°C for 10 min. Modified PCR programs, including touchdown PCR, were used to optimize some primer pairs when necessary. Products were pooled post-PCR and resolved on an ABI 377XL sequencer using GS-500 ROX (Applied Biosystems) internal size standard in each lane. Gel image data were processed and alleles scored using GeneScan version 3.1 and Genotyper version 2.1 software (Applied Biosystems).

### Chromosome Maps

SSLP markers were selected from the Dietrich et al. (1996) set. Map positions were obtained from the Mouse Genome Database (TJL). Graphical linkage maps for each chromosome was generated using Encyclopedia of the Mouse Genome 3.0, Version 1.0a17 for the Macintosh (TJL).

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- <http://www.cidr.jhmi.edu/mouse/mouse.html>; CIDR-Mouse  
Genotyping Home Page.
- <http://www.informatics.jax.org>; Mouse Genome Database Home  
Page.
- <http://www.genome.wi.mit.edu/cgi-bin/mouse/index#genetic>;  
WICGR Resources, Genetic and Physical Maps of the Mouse  
Genome.
- [http://www.ensembl.org/Mus\\_musculus/](http://www.ensembl.org/Mus_musculus/); Ensembl Genome Browser,  
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**Systematic Discovery of New Genes in the *Saccharomyces cerevisiae* Genome**

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Tira M. Templeton's name was spelled incorrectly in the above article. It should be Tiva M. Templeton.





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