

## *Prdm9* Controls Activation of Mammalian Recombination Hotspots

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Genetic recombination is an essential biological process among eukaryotes. Mammalian meiotic recombination, which preferentially occurs at specialized sites, 1-2 Kb long, known as hotspots, assures the orderly segregation of meiotic chromosomes and creates genetic variation among offspring. However, despite their importance in determining the recombination landscape of the organism, we have little understanding of the elements determining the location and relative activity of hotspots. That the location of a hotspot is not determined simply by its internal DNA sequence is supported by both yeast (1) and mammalian studies (2,3). Two recent reports (4,5) have described the existence of trans-acting loci that control the activation of specific hotspots elsewhere. The loci, *Dsbc1* and *Rcr1*, are located in overlapping 5.4 Mb and 6.3 Mb regions on mouse chromosome 17.

We have now extended the mapping of both *Rcr1* and *Dsbc1* using 1580 progeny of a cross between the C57BL/6J (B6) and CAST/EiJ (CAST) mouse strains differing in activity of the *Rcr1/Dsbc1* controlled hotspots *Hlx1*, *Esrrg-1* and *Psmb9* (fig. S1). We located both loci to a common 181 Kb region containing four protein coding genes (Fig. 1): *Psmbl1*, *Tbp*, *Pdcd2*, and *Prdm9*. We found no differences in expression levels of the transcripts of any of these proteins in testes when assayed by RT-PCR (table S1).

*Psmbl1*, proteasome beta type 1 subunit, is a ubiquitously expressed protein that functions as a structural component of an organelle responsible for generalized protein degradation. *Tbp*, TATA binding protein, is an essential component of the TFIID transcription initiation complex which has considerable DNA binding specificity. *Pdcd2*, programmed cell death protein 2, is widely expressed, repressed during B cell lymphomagenesis. Based on DNA sequencing of the entire coding regions of these three proteins, we found only five coding SNPs, all synonymous, resulting in no amino acid differences between the B6 and CAST strains (table S2). For the first exon of *Pdcd2*, we relied on the Sanger Institute sequence (<http://www.sanger.ac.uk/modelorgs/mousegenomes/>).

*Prdm9*, PR domain containing 9, is uniquely expressed during early meiosis in both males and females, and the knockout is blocked at the pachytene stage of meiosis I with reduction of the number of *Dmcl1* foci and a loss of *Dmcl1* and  $\gamma$ -*H2AX* colocalization, resulting in sterility in both sexes and azoospermia in males (6). It has three functional domains, a N terminal KRAB domain that can promote protein:protein binding and transcriptional repression when tethered to DNA by an adjacent DNA binding domain, a central PR/SET

domain providing a histone methyl transferase activity capable of trimethylating H3K4 and thus altering chromatin configuration, and a terminal zinc finger domain of C2H2 type. The zinc finger domain contains tandem repeats, with one finger per repeat, and has 11 fingers in the CAST allele and 12 in the B6 allele, with a number of amino acid substitutions in the 6, 9 and 12 positions after the second cysteine of the fingers, the residues determining DNA binding specificity. (fig. S2). Given the properties of *Prdm9*, including its requirement for meiosis, its histone 3 lysine-4 trimethylation activity and role in determining DNA DSBs, and the lack of a reasonable alternative candidate, we conclude that *Prdm9* is *Rcr1/Dsbc1*. This is supported by the observation (7) that recombination at hotspots *Psmb9* and *Hlx1*, regulated by *Rcr1/Dsbc1*, is preceded by histone 3 lysine-4 trimethylation.

We sequenced *Prdm9* exon 12 containing the entire Zn finger domain in 20 mouse strains and found five alleles differing in the number of zinc finger repeats, ranging from 11 to 14, as well as in codons for the amino acids in 6<sup>th</sup>, 9<sup>th</sup> and 12<sup>th</sup> position of each finger responsible for DNA binding (fig. S2). The alleles for C3H and PWD mouse strains are identical with those reported by Mihola et al. (8) who identified *Prdm9* as the gene underlying the *Hst1* locus determining male infertility in certain crosses between strains of *M.m. musculus* and *M.m. domesticus*.

We also sequenced exon 12 of human *PRDM9* in 64 DNA samples from the Coriell Institute for Medical Research. We found two predominant alleles among these 128 chromosomes, containing a zinc finger domain of 12 tandem repeats, and several minor frequency alleles (fig. S3A). African-Americans had eleven minor alleles with 11, 12 or 13 zinc fingers and a number of amino acid substitutions, Han Chinese three minor alleles, two with 12 and one with 13 fingers, and Mexican-Americans one, identical to the Chinese variant with 13 fingers, while Caucasians had none (fig. S3B). These results conform well with presently accepted views of human evolution.

The discovery of *Prdm9* as the first mammalian protein regulating meiotic recombination hotspots initiates studies of an important biological control system that has hitherto been inaccessible. A recent paper (9) identified *Prdm9* as a speciation gene across diverse metazoans and hypothesized that its essential role in meiosis is directly related to its ability to bind rapidly evolving DNA sequences. Our results clearly show that these sequences represent recombination hotspots. An immediate compelling question is whether *Prdm9* controls activation of all, or nearly all, recombination

hotspots, or whether it is simply a member of a family of proteins, each controlling a sub-set of all hotspots.

order and coordinates of the four protein-coding genes in the critical region.

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## Supporting Online Material

[www.sciencemag.org/cgi/content/full/science.1181495/DC1](http://www.sciencemag.org/cgi/content/full/science.1181495/DC1)

Materials and Methods

Figs. S1 to S3

Table S1

References

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**Fig. 1.** Fine mapping the location of *Rcr1*. Upper line shows the position, in Mb (Ensembl Build 37), of the markers used for genetic mapping. Crossovers collected in this region are marked with blue for B6 DNA and red for CAST. On the left are the serial numbers of the male progeny tested for recombination activity. The critical region containing *Rcr1* is between markers at 15.589 and 15.770 Mb. Below are the

