Summary of publications

Positional cloning of the Hybrid sterility 1 gene: fine genetic mapping and evaluation of two candidate genes

Hybrid sterility is one of the mechanisms of speciation. The Hybrid sterility 1 (Hst1) gene was the first mapped mammalian gene. The gene affects fertility of male hybrids between certain laboratory strains (such as C57BI/10) and Mus musculus musculus mice by causing a breakdown of spermatogenesis at the stage of primary spermatocytes. In the process of positional cloning of the Hst1 gene, we generated a contig of bacterial artificial chromosomes (BACs) and subsequently a low coverage sequence of the candidate region of the 129S1/SvImJ strain. New genetic markers narrowed down the Hst1 region from 580 to 360 kilobases. The products of two genes from this region, TATA-binding protein (Tbp) and proteasome subunit beta 1 (Psmb1), accumulate during spermatogenesis. The proteins have been described previously as having conserved C-terminal sequences and species-specific Ntermini. We evaluated the candidacy of these genes for Hst1 by allelic sequencing and by real-time reverse-transcription PCR of testicular mRNAs. The results suggest that neither the Psmb1 nor the Tbp gene cause hybrid sterility. The single nucleotide polymorphisms (SNPs) we have found, was used for the haplotype analysis of the Hst1 region.

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