

Research

A recombination hotspot leads to sequence variability within a novel gene (*AK005651*) and contributes to type 1 diabetes susceptibility

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More than 25 loci have been linked to type 1 diabetes (T1D) in the nonobese diabetic (NOD) mouse, but identification of the underlying genes remains challenging. We describe here the positional cloning of a T1D susceptibility locus, *Idd11*, located on mouse chromosome 4. Sequence analysis of a series of congenic NOD mouse strains over a critical 6.9-kb interval in these mice and in 25 inbred strains identified several haplotypes, including a unique NOD haplotype, associated with varying levels of T1D susceptibility. Haplotype diversity within this interval between congenic NOD mouse strains was due to a recombination hotspot that generated four crossover breakpoints, including one with a complex conversion tract. The *Idd11* haplotype and recombination hotspot are located within a predicted gene of unknown function, which exhibits decreased expression in relevant tissues of NOD mice. Notably, it was the recombination hotspot that aided our mapping of *Idd11* and confirms that recombination hotspots can create genetic variation affecting a common polygenic disease. This finding has implications for human genetic association studies, which may be affected by the approximately 33,000 estimated hotspots in the genome.

[Supplemental material is available online at <http://www.genome.org>. The sequence data from this study have been submitted to dbSNP (<http://www.ncbi.nlm.nih.gov/projects/SNP/>) under accession nos. ss262803370, ss262803372, ss262803374, ss262803376, ss262803379, ss262803382, ss262803385, ss262803388, ss262803390, ss262803391, ss262803392, ss262803394, ss262803397, ss262803400, ss262803402, ss262803403, ss262803404, and ss262803405, and to the NCBI Probe Database (<http://www.ncbi.nlm.nih.gov/sites/entrez?db=probe>) under accession nos. 10544425–10544446.]

Type 1 diabetes (T1D) is a polygenic autoimmune disease in which lymphocytes mediate the destruction of insulin-producing beta cells in the pancreas (Atkinson and Eisenbarth 2001). The events that trigger the pathogenic autoimmune response are still not clear, but recent genome-wide association studies indicate that there are more than 40 loci affecting susceptibility to T1D in humans (Hakonarson et al. 2007; Todd et al. 2007; The Wellcome Trust Case Control Consortium 2007; Barrett et al. 2009; Concannon et al. 2009a). These studies have detected association of T1D with the common variants for previously identified genes and new candidates. Except for the HLA locus, however, these loci have small effects upon disease risk (odds ratio < 2.5) and fail to adequately explain the genetic variance for T1D (Concannon et al. 2009b). Instead, it has been proposed that rare/private mutations with larger effects may account for the missing genetic variance in complex genetic diseases (Goldstein 2009). Whether rare or

common, identifying the actual causative variants for T1D has proved challenging due to genetic heterogeneity among affected individuals.

A complementary approach to human studies is the use of inbred mouse strains, in which genetic heterogeneity is avoided, and selective mating can precisely map genes for which allelic variation affects disease susceptibility. Analysis of the nonobese diabetic (NOD) mouse strain, which spontaneously develops T1D similar to humans, has been widely used to better understand disease pathology and gain key insights into the genetics of T1D (Atkinson and Leiter 1999). In parallel to human studies, more than 25 loci (termed *Idd*) have been linked to T1D in the NOD mouse (Serreze and Leiter 2001; Ridgway et al. 2008). Confirmation of these loci is best achieved using congenic mouse strains (Rogner and Avner 2003), which are generated by controlled mating of NOD mice with diabetes-resistant strains to introduce a donor-derived chromosome interval carrying a resistant allele onto the susceptible NOD genetic background. By testing smaller donor-derived intervals for their effect upon diabetes onset, a region small enough to be sequenced for disease-causing variants can be identified. To date, congenic NOD strains have confirmed *Idd* loci on chromosomes (chr) 1–4, 6, 7, 11, 13, 17, and 18 (Serreze and

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Leiter 2001; Ridgway et al. 2008). Besides the MHC region on Chr 17, at least eight of these loci have been dissected into smaller intervals using congenic NOD strains, and accumulating evidence has identified *B2m*, *Il2*, *Il21*, *Ctla4*, *Slc11a1*, and *Tpvt1* as genes for which NOD mice harbor T1D susceptibility alleles (Hamilton-Williams et al. 2001; Kissler et al. 2006; Razavi et al. 2006; Yamanouchi et al. 2007; Araki et al. 2009; McGuire et al. 2009). Preliminary evidence also suggests that *Arntl2* and *Stat5b* are potential T1D susceptibility genes (Hung et al. 2006; Laloraya et al. 2006).

Although few causative variants have been defined to date, the NOD alleles that increase T1D risk in this mouse strain can be either rare or common among inbred mouse strains. For example, the effect of the *Idd1* locus is attributed to the relatively rare MHC class II variant (*Ab^{g7}*), together with a more common variant found in other inbred mouse strains that encodes a deletion in the I-E- α chain promoter (Serreze and Leiter 2001). This observation, along with human genetic studies, suggests that increased T1D risk in humans may also result from the combination of rare and common variants within the human population (Concannon et al. 2009b).

Despite the identification of several *Idd* genes to date, this limited collection does not fully explain T1D pathogenesis or the underlying genetic architecture for T1D risk. One of the many *Idd* loci still to be identified is *Idd11*, which is located on Chr 4 and originally linked to T1D in NOD backcrosses to the C57BL/6 (B6) and SJL strains (Morahan et al. 1994). As B6 mice carry a resistance allele for *Idd11*, congenic strains with different Chr 4-B6-derived intervals on the NOD genetic background were produced (NOD.B6Idd11A, NOD.B6Idd11B, NOD.B6Idd11C, NOD.B6Idd11D). Three of these congenic NOD strains demonstrated significant diabetes resistance, thus confirming and localizing *Idd11* to an ~8-Mb interval on Chr 4 (Brodnicki et al. 2000, 2005).

To localize *Idd11* further and identify the underlying gene, we established new congenic NOD strains to dissect this ~8-Mb interval and monitored them for diabetes onset. Remarkably, each of these smaller congenic intervals were derived from recombination breakpoints within the same 6.9-kb interval that resulted in varying levels of T1D susceptibility for the different congenic NOD strains. Here, we report the sequence analysis of these breakpoints and the identification of a recombination hotspot that led to the discovery of a novel candidate gene (GenBank mRNA: *AK005651*) for *Idd11*. This gene of unknown function exhibits decreased expression in the thymus and spleen of NOD mice. Furthermore, NOD mice carry a unique haplotype for *Idd11* compared with 25 other inbred mouse strains analyzed. Our findings demonstrate that recombination hotspots, which have been relatively neglected in human association studies, can create unique DNA sequence variation that has relatively large effects upon the risk for a common polygenic disease.

Results

Mapping *Idd11* using congenic mouse strains

To localize *Idd11* further, new congenic mouse strains were derived from NOD.B6Idd11D because this strain carried the smallest Chr 4-B6-derived interval providing diabetes protection among our panel of previously characterized congenic strains (Table 1; Brodnicki et al. 2000, 2005). Briefly, heterozygous NOD.B6Idd11D mice were intercrossed to generate F₂ progeny that were screened for recombination events by genotyping novel markers we identified within this interval (Table 1; Supplemental Table 1; Supplemental Fig. 1). Three new congenic NOD strains (NOD.B6Idd11E, NOD.B6Idd11F,

Table 1. Genetic intervals for *Idd11* congenic mouse strains

Marker ^a	~Mb ^b	Congenic strains ^c					
		B ^d	D	E	F	G	
<i>D4Mit12</i>	124,048,407	N	N	N	N	N	
<i>D4Mit338</i>	125,017,654	N	N	N	N	N	
<i>D4Mit73</i>	126,497,935	N	N	N	N	N	
<i>D4Mit72</i>	128,630,230	N	B	B	N	B	
<i>D4Mit203</i>	129,249,262	N	B	B	N	B	
<i>D4Wehi1</i>	129,422,665	N	B	B	N	B	
<i>D4Wehi2</i>	129,521,871	N	B	B	N	B	
<i>D4Wehi5</i>	129,633,461	N	B	B	N	B	
		*	B	*	*	*	<i>Idd11</i>
<i>D4Wehi6</i>	129,640,320	B	B	N	B	N	
<i>D4Wehi3</i>	129,666,984	B	B	N	B	N	
<i>D4Wehi17</i>	129,711,291	B	B	N	B	N	
<i>D4Wehi21</i>	130,392,752	B	B	N	B	N	
<i>A892</i>	130,848,378	B	B	N	B	N	
<i>D4Wehi22</i>	132,008,637	B	B	N	B	N	
<i>D4Mit204</i>	132,983,282	B	B	N	B	N	
<i>D4Mit339</i>	133,923,341	B	B	N	B	N	
<i>D4Mit69</i>	135,916,989	N	N	N	N	N	
<i>D4Mit126</i>	142,152,658	N	N	N	N	N	
<i>D4Mit256</i>	154,364,548	N	N	N	N	N	

^aFor *D4Wehi* marker oligonucleotides and their NCBI Probe Database accession numbers, see Supplemental Table 1.

^bGenomic coordinates are from NCBI build 37 assembly, mm9.

^cStrain names have been abbreviated (e.g., D = Idd11D = NOD.B6Idd11D).

^dThe Idd11B congenic interval is presented here for comparison. The T1D incidence curve for Idd11B mice has been previously reported and is similar to NOD mice (Brodnicki et al. 2000, 2005).

B (boldface), C57BL/6 genotype; N, NOD genotype; *, location of breakpoints described in Table 2 that aided in localizing *Idd11*.

NOD.B6Idd11G) were established from recombinant F₂ mice, and these were monitored for diabetes onset compared to NOD and Idd11D mice (note that henceforth congenic strain names are abbreviated, e.g., NOD.B6Idd11D = Idd11D) (Fig. 1).

Comparison of the resulting diabetes incidence curves indicated that *Idd11* mapped to an ~6.9-kb interval, between *D4Wehi5* and *D4Wehi6* (Table 1; Fig. 1A,B). However, the *Idd11* locus appeared to be more complex than expected because the congenic strains exhibited varying levels of T1D incidence. For example, Idd11E and Idd11G seemed to have identical B6-derived intervals, but Idd11E was more susceptible to T1D than Idd11G (65% vs. 33% diabetic by 300 d). On the other hand, the Idd11G interval provided less diabetes protection compared with the larger Idd11D interval (33% vs. 6%) but provided greater protection compared with the Idd11F interval (33% vs. 52%). As Idd11E, Idd11F, and Idd11G were derived from the Idd11D strain, we postulated that sequencing the recombinant boundary between *D4Wehi5* and *D4Wehi6* would explain the variability in T1D susceptibility between the congenic strains.

Sequence analysis of the *Idd11* critical interval

Sequence analysis of the NOD, B6, and congenic NOD strains identified several sequence variants within the ~6.9-kb interval (Table 2; Supplemental Tables 2, 3). Remarkably, Idd11E and Idd11G were isogenic except at sequence variant 3, demonstrating that allelic variation at this position can significantly affect T1D susceptibility. However, the B6-derived allele at this position alone could not account for all of the *Idd11* effect since it conferred different levels of diabetes protection to Idd11D (~6% diabetic by 300 d), Idd11F

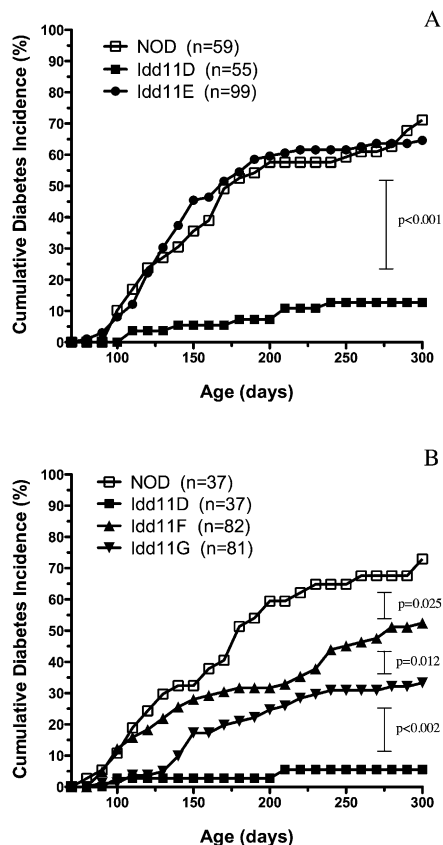


Figure 1. Cumulative diabetes incidence curves for *Idd11* congenic strains. Female cohorts were monitored for diabetes by measuring urinary glucose levels. Pairwise comparisons of diabetes incidence curves were performed using the log-rank test. As a historical footnote, Idd11E was monitored first because it was presumed that this strain would be protected against diabetes given that Idd11B exhibited a similar diabetes incidence curve to NOD (Brodnicki et al. 2000). Once we observed that Idd11E was not protected against diabetes, we established cohorts for Idd11F and Idd11G to confirm this result using independent congenic strains, which initially appeared to have congenic intervals similar to Idd11B and Idd11E, respectively (Table 1). Subsequent sequence analysis of the interval between *D4Wehi5* and *D4Wehi6* (Table 2) identified genetic variation that explained the diabetes incidence observed for Idd11B (Brodnicki et al. 2000) and Idd11E (A).

(~52%), and Idd11G (~33%) (Fig. 1A,B; Table 2). This comparison of diabetes incidence curves indicated that *Idd11* was likely due to a haplotype effect in which B6-derived sequences are required at more than one position to provide optimal T1D protection.

These data raised a critical question: How big is the *Idd11* haplotype? NOD crosses with either SJL, NON, 129, C57L, C57BL/10 (B10), or NOR mouse strains have previously demonstrated the linkage of T1D to chr 4, suggesting that these strains, similar to B6, may harbor a T1D-resistance allele for *Idd11* (Morahan et al. 1994; Rodrigues et al. 1994; McAleer et al. 1995; McDuffie 2000; Reifsnnyder et al. 2005; Leiter et al. 2009). The B10-identified and the NOR-identified loci, which overlap the B6-defined *Idd11* locus, have also been confirmed by congenic strains (Lyons et al. 2000; Reifsnnyder et al. 2005). Sequence analysis determined that the B10 and NOR strains were identical to B6 across the congenic breakpoint positions (Table 2; Supplemental Table 4). In contrast, the SJL strain was identical to NOD at these variants except at 1, 3, and 5, for which SJL was identical to B6, B10, and NOR (Table 2; Supplemental Table

4). Further sequence analysis for these variants within 20 other inbred mouse strains (including wild-derived strains) indicated that the *Idd11* haplotype consisted of at least variants 1–5, with NOD mice representing a unique haplotype marked by a 12-bp deletion at variant 1 (Table 2; Supplemental Table 4).

Confirmation of a recombination hotspot

It was conspicuous that recombination breakpoints for four of our congenic strains occurred within an ~1.2-kb interval and resulted in one instance of a crossover with a complex conversion tract (Idd11E) (Table 2). This dense clustering of crossover events and the presence of a complex conversion tract is characteristic of a recombination hotspot (Petes 2001; Jeffreys and May 2004; Bois 2007; Kauppi et al. 2007). To confirm and measure the frequency of the meiotic crossovers within the *Idd11* haplotype, 723 F₂ progeny were generated by intercrossing heterozygous (NOD × Idd11D)F₁ mice and were screened for recombination events between *D4Wehi5* and *D4Wehi6*. The calculated crossover activity was ~50 cM Mb⁻¹, which is 100-fold greater than the mouse genome average (~0.5 cM Mb⁻¹) (Shiroishi et al. 1995), and confirms that this interval harbors a recombination hotspot (Fig. 2). Genomic DNA available for 231 of these F₂ progeny (representing 462 meiosis events) that did not exhibit a crossover between *D4Wehi5* and *D4Wehi6* was further genotyped for sequence variants 1–8 (Table 2). Only one noncrossover event (also termed gene conversion) was identified, at variant 4. This suggests a relatively low frequency (<0.5%) for noncrossover events in this hotspot, but gene conversion events can only be detected and estimated using the available sequence variants located within the hotspot interval. Hence, this conversion frequency is likely underestimated due to the relatively few variants within this interval between NOD and B6 mice. While other local recombination hotspots may reside elsewhere within the Idd11D interval, we observed no recombination events either ≤112 kb proximal to *D4Wehi5* or ≤71 kb distal to *D4Wehi6* (Fig. 2, inset). These flanking recombination “cold regions” have restricted our ability to generate smaller congenic intervals encompassing only the proposed *Idd11* haplotype. It should be noted that while this recombination hotspot may generate noncrossover events, more than one variant within this haplotype would need to be converted to confer T1D protection in subsequent congenic NOD mice.

Bioinformatic and expression analysis of a novel candidate gene for *Idd11*

The *Idd11* haplotype, including the recombination hotspot, is located within a predicted gene, termed *AK005651* (i.e., GenBank mRNA accession number), supported by a collection of expressed sequence tags (ESTs) (Supplemental Fig. 2). The encoded transcript consists of five exons (Fig. 3A; Supplemental Table 5) and two splice isoforms (±exon 3) confirmed by RT-PCR and sequencing data (data not shown). *AK005651* has a number of open reading frames, but none demonstrate clear protein domains or homology with known proteins nor does the transcript encode an obvious microRNA. Conservation between mammalian sequences across this region suggests the presence of a human ortholog, although no orthologous human spliced EST or transcript has yet been described (bioinformatic analysis is summarized in Supplemental Table 6).

If *AK005651* does encode a protein, then variants 2 and 4 result in amino acid changes that depend on the open reading

Table 2. Genotypes for the proposed *Idd11* haplotype

	Sequence variant ^a								Cumulative diabetes incidence
	1	2	3	4	5	6	7	8	
NOD	—	C	C	A	T	C	—	T	71%, 73%
Idd11B ^b	—	C	C	A	C	T	AG	C	73%
Idd11E ^c	CCTCTCGGTGTT	T	C	G	T	C	—	T	65% ^c
Idd11F	—	T	T	G	C	T	AG	C	52%
Idd11G	CCTCTCGGTGTT	T	T	G	T	C	—	T	33%
Idd11D	CCTCTCGGTGTT	T	T	G	C	T	AG	C	6%, 12%
NON ^d	CCTCTCGGTGTT	C	C	A	C	C	—	T	0%
SJL ^e	CCTCTCGGTGTT	C	T	A	C	C	—	T	0%
PWD	CCTCTCGGTGTT	C	T	G	C	C	AG	C	0% ^g
CAST	CCTCTCGGTGTT	T	T	G	C	C	AG	T	0% ^g
C57BL/6 ^f	CCTCTCGGTGTT	T	T	G	C	T	AG	C	0%
Distance between markers (bp)		128	138	149	37				
			152	820	557				

^aSequence variant 1, ss262803379 (129,638,063); 2, rs32465505 (129,638,192); 3, rs27507073 (129,638,344); 4, rs27507071 (129,638,482); 5, rs27507066 (129,639,302); 6, rs27507065 (129,639,451); 7, ss262803405 (129,640,008); and 8, rs48826903 (129,640,045). Chr 4 positions are listed in parentheses and based on NCBI build 37 assembly, mm9. Boldface text indicates sequence identical to C57BL/6 sequence.

^bIdd11B mice exhibit a similar diabetes incidence curve to NOD and Idd11E mice (Brodnicki et al. 2000, 2005).

^cIdd11E represents a crossover with a complex conversion tract as confirmed by direct sequencing of a PCR product encompassing markers 1–4 derived from genomic DNA from five independent Idd11E mice.

^dVariants shared by NON, CTS, BALB/c, NZO, NZW, C3H, CBA, DBA/2, A/J, ALR, ALS, AKR, 129/Sv, WSB.

^eVariants shared by SJL, SWR, FVB.

^fVariants shared by C57BL/6, C57BL/10, NOR, NZB, DBA/1, MOLF.

^gSpontaneous diabetes has not been reported for PWD and CAST mouse strains. Note that sequencing of genomic DNA obtained from independent NOD and B6 strains from the Jackson Laboratory were identical to the NOD and B6 strains within our mouse colony. Sequence variation across this region showing the individual mouse strains is provided in Supplemental Table 4.

frame (as shown in Supplemental Table 7). For the three possible open reading frames, only one results in silent substitutions due to variants 2 and 4 (Supplemental Table 7, ORF 1), whereas these two variants produce amino acid changes in all other open reading frames between NOD and B6. The longest open reading frame for *AK005651* (methionine to stop) is 53 amino acids and is located in exon 5. No sequence variation between NOD and B6 occurs in this open reading frame.

Sequence variation within the proposed *Idd11* haplotype may also affect gene expression. The sequence encompassing variants 1–4 contains predicted transcription factor binding sites (TFBSs) based upon conservative analyses using TRANSFAC matrices (Matys et al. 2006), but none of the *Idd11* sequence variants occur within these predicted TFBSs (Supplemental Table 8). Variants 1 and 2 do occur in a promoter-associated regulatory feature (Supplemental Fig. 2) predicted by Ensembl based upon DNase I hypersensitivity and the enrichment of histone H3K4me3 in different cell types (Mikkelsen et al. 2007). Real-time PCR analysis indicated that *AK005651* is differentially expressed between NOD and diabetes-resistant mouse strains (i.e., B6 and Idd11D) in tissues relevant to T1D pathogenesis. Exon 4 and 5 were present in both splice variants, and B6 and Idd11D mice had at least a twofold increase in thymic and splenic expression of these spliced exons (Fig. 3B). On the other hand, expression analysis of *Spocd1* and *Bai2*, two genes flanking *AK005651* (Supplemental Fig. 2), indicated that the sequence variation within *AK005651*, in particular variants 1–4 for the *Idd11* haplotype, is unlikely to affect transcription levels of these two flanking genes (Supplemental Fig. 3).

Expression differences may also reflect deficient exon splicing. Sequence variants 2–5 do not disrupt canonical splice sites,

although they may alter cryptic intron/exon splicing motifs (Chasin 2007; Wang and Burge 2008). In contrast, the unique NOD 12-bp deletion at position 1 shortens the intron between exons 2 and 3. Real-time PCR detected a significant decrease in a spliced product from these two exons in thymic and splenic RNA isolated from NOD mice compared with B6 and Idd11D mice (Fig. 3C). The spliced product from exons 2 and 4 was also significantly decreased in the NOD thymus and spleen, as well as the pancreas, compared with B6 and Idd11D (Fig. 3D). We note that the anomalous result for the liver (Fig. 3C) was due to one extreme outlier. This B6 sample did not result in an outlier using other real-time PCR probes and was the only such outlier observed within our expression data set (Figs. 3, 4; Supplemental Fig. 3).

To further dissect the effect of the *Idd11* haplotype upon *AK005651*, expression analysis was performed using thymic and splenic RNA isolated from our series of congenic mouse strains (Fig. 4; note: NOD, B6, Idd11D represent different cohorts to those in Fig. 3). Variant 1 (12-bp ins/del) had the most distinct effect upon *AK005651* expression. Congenic mice harboring the B6-derived insertion (Idd11D, Idd11G, Idd11E) ex-

hibited significantly higher transcription levels of the spliced product for exons 2/3 compared with NOD mice (Fig. 4B). However, the effect of the *Idd11* haplotype upon the transcription levels of spliced products for exons 2/4 and for exons 4/5 appears to be more complex. While the two spliced products had similar expression profiles in the respective tissues (Fig. 4A,C), the different *Idd11* haplotypes led to varied expression between strains, as well as between tissues. For example, only Idd11D exhibited twofold or more increases for both splice products in both tissues; whereas Idd11E and Idd11F had more than twofold increases only in the spleen. These results indicate that different combinations of B6-derived sequences within the *Idd11* haplotype have varying effects upon *AK005651* expression, with increased expression for all three spliced products in both the thymus and spleen correlating with the greatest degree of T1D protection (i.e., Idd11D).

Discussion

Positional cloning using congenic mouse strains has been used to confirm and localize a number of T1D susceptibility loci (Serreze and Leiter 2001; Ridgway et al. 2008). Nonetheless, only a handful of genes for *Idd* loci have been identified to date, principally because it takes large mouse colonies and significant breeding time to generate a series of congenic mouse strains that maps a locus to within a genomic interval containing only one or a few candidate genes. Even when the candidate genes are identified (whether in mouse or human genetic studies), it is often difficult to determine which sequence variant(s) within the mapped interval are causal, unless they are obvious mutations that disrupt known coding, splicing, or regulatory motifs (Hindorf et al. 2009). In this study,

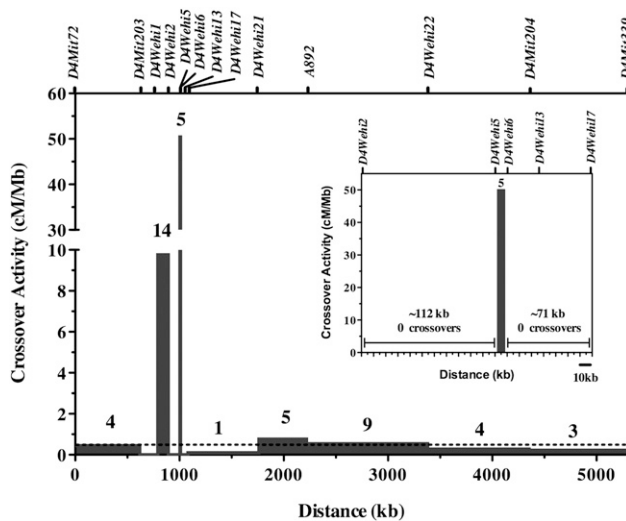


Figure 2. Distribution of crossover breakpoints within the NOD.B6Idd11D congenic interval. Seven-hundred-twenty-three F_2 pups (1446 meiosis events) were derived from intercrossing NOD.B6Idd11D F_1 mice and were genotyped for markers within the NOD.B6Idd11D congenic interval. The crossover activity was calculated by dividing the frequency of crossovers in each interval by its length. For example, five pups had a recombination between *D4Wehi5* and *D4Wehi6*, giving rise to a genetic distance of 0.345 cM across this 6859-bp interval, which is ~ 50 cM Mb $^{-1}$. Bars, crossover frequency in cM/Mb in each interval. Above each bar, the number of crossovers observed in that marker interval is shown. The dashed line indicates the average crossover activity for the mouse genome: ~ 0.5 cM/Mb (Shiroishi et al. 1995). Genetic markers are indicated by ticks above the plot. The inset displays an enlarged view of the interval between *D4Wehi2* and *D4Wehi7*.

a recombination hotspot generated a unique series of congenic mouse strains for *Idd11* and enabled us to identify a novel candidate gene for which sequence variation affects T1D susceptibility in the NOD mouse.

Comparison of the sequence and diabetes incidence curves between our congenic NOD strains indicated that *Idd11* is due to a haplotype, consisting of at least five variants, for which the composition has variable effects on susceptibility to T1D. For example, the Idd11D congenic strain has the complete B6-derived haplotype and the best protection against diabetes onset, whereas Idd11E and Idd11B lack two or more B6-derived alleles within this haplotype and have similar diabetes incidence to NOD mice (Table 2; Fig. 1; Brodnicki et al. 2000). In contrast, Idd11F and Idd11G mice lack one B6-derived allele within the proposed *Idd11* haplotype but do so at different positions and exhibited different T1D incidence curves (Table 2; Fig. 1). Thus these variable effects depend on the number and combination of B6-derived alleles within this haplotype. Similar associations of polymorphic haplotypes with autoimmune disease are well established for the major histocompatibility complex (Fernando et al. 2008), the human cytokine gene cluster on chr 5q31 (Rioux et al. 2001), and the mouse SLAM/CD2 gene cluster (Wandstrat et al. 2004). However, the *Idd11* haplotype appears to be confined to a single gene, whereas these other haplotypes encompass multiple genes.

Sequence analysis determined that NOD mice have a unique *Idd11* haplotype due to a 12-bp deletion. Notably, the NOD strain was generated from the same inbreeding program as the CTS and NON strains (Makino et al. 1980; Beck et al. 2000), but these two strains (as well as 18 other laboratory and four wild-derived inbred

strains) do not have the 12-bp deletion, suggesting this microdeletion arose as a de novo mutation or was preferentially inherited from a different parental ancestor during the derivation of the NOD strain. In either case, further sequence comparison was required because this microdeletion did not fully explain the varying levels of T1D susceptibility observed in our congenic mouse strains. Two additional sequence variants (rs27507073 and rs27507066) were associated with T1D based on resistance phenotypes determined by segregation and congenic analyses (Brodnicki et al. 2000, 2005; Lyons et al. 2000; Reifsnnyder et al. 2005). Our finding does not imply that the *Idd11* haplotype is limited to these three variants, but only indicates the minimum size of the B6-derived haplotype providing protection against T1D in NOD mice.

Idd11 is not the only T1D susceptibility locus on chr 4. Genetic studies have mapped at least three other *Idd* loci to this chromosome. An outcross between NOD and B10 originally detected a locus, termed *Idd9*, linked next to the telomere of chr 4 (Rodrigues et al. 1994). Congenic NOD strains, harboring B10-derived intervals, subsequently expanded and dissected this locus into three subloci (*Idd9.1*, *Idd9.2*, *Idd9.3*) (Lyons et al. 2000), with *Idd9.1* mapped to the same region we had previously localized *Idd11* (Morahan et al. 1994). The haplotype analysis reported here suggests that *Idd11* and *Idd9.1* are the same because B6 and B10 are identical by descent for this chromosome region. A NOD outcross with the NOR strain also detected linkage to the region encompassing *Idd11*, which was confirmed by NOD.NOR-chr 4 congenic mice (Reifsnnyder et al. 2005). NOR is an inbred recombinant congenic strain derived from NOD and C57BLKS/J (Prochazka et al. 1992), and our sequence analysis indicated that the *Idd11* haplotype for NOR is B6-derived. However, NOD.NOR-chr 4 mice harbor a larger congenic interval than the B6-derived intervals in our *Idd11* congenic strains. Thus, resistant NOR-derived alleles at other chr 4 loci may also contribute to the overall diabetes protection observed for this congenic strain (Reifsnnyder et al. 2005). Lastly, NOD outcrosses with the NON and 129/SvImJ mouse strains also detected linkage of T1D to chr 4 (McAlee et al. 1995; Leiter et al. 2009), but it is more likely that these linkage results were due to allelic variation at other loci because these two strains are NOD-like for the *Idd11* haplotype—their allelic composition is not predicted to be protective based on comparison with the Idd11E haplotype for *Idd11* (Table 2).

Idd11 appears to represent a “gene-based functional haplotype”: a defined sequence interval taken as a unit because individual variants are not sufficient to act as separate disease markers or completely account for the associated phenotypic effect (Hoehe 2003). The *Idd11* haplotype, including the recombination hotspot, is located within *AK005651*, a predicted gene of unknown function. None of the open reading frames demonstrate homology with known proteins or evolutionary conservation, suggesting that *AK005651* may encode a long noncoding RNA (Mercer et al. 2009; Ponting et al. 2009), although it was not detected in a recent large noncoding RNA screen of four mouse cell types (Guttman et al. 2009). As current bioinformatics approaches are limited in deciphering the function of *AK005651*, we investigated the effect of sequence variation on *AK005651* expression.

The different *Idd11* haplotypes, represented by our series of congenic strains, have variable effects on *AK005651* expression, which is perhaps not surprising given their variable effects on susceptibility to T1D. Real-time PCR indicated that the NOD *Idd11* haplotype is associated with decreased expression of *AK005651* in the thymus and spleen. In particular, the unique NOD microdeletion was associated with the decreased expression of the exon

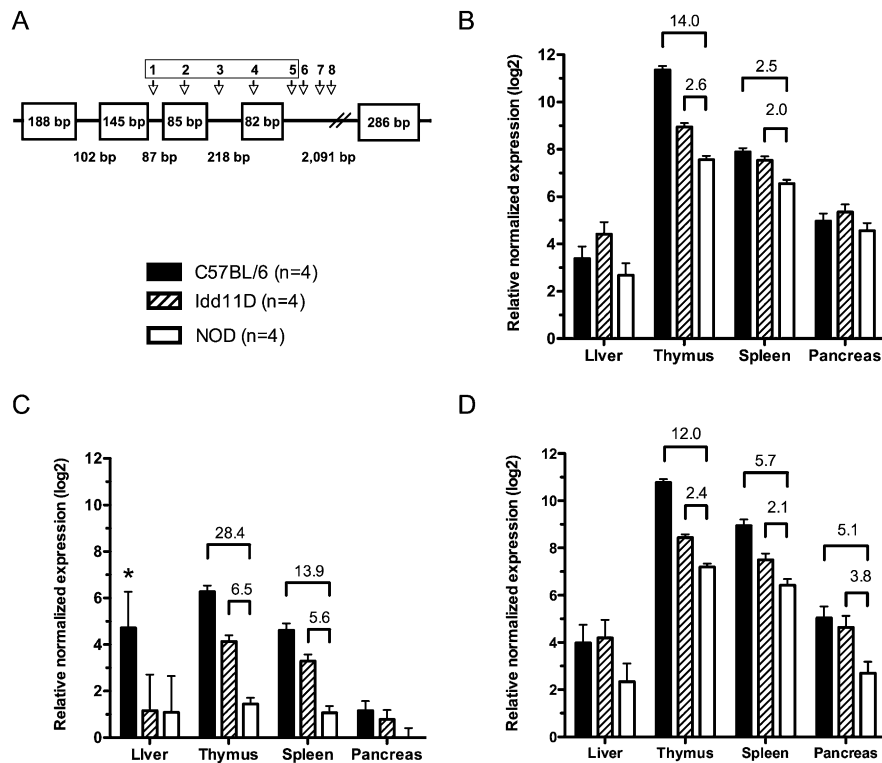


Figure 3. Schematic diagram and expression analysis of *AK005651*. (A) The *Idd11* haplotype (variants 1–5) is located within a predicted gene, termed *AK005651*, consisting of five exons (Supplemental Table 5). Exon 3 is alternatively spliced giving rise to two different transcripts (\pm exon 3). Quantitative real-time PCR was performed to detect expression differences between mouse strains at 50 d of age for the spliced product derived from: (B) exons 4 and 5, which are present in both splice variants; (C) exons 2 and 3, (*) one of the four C57BL/6 samples was an extreme outlier (the mean relative normalized expression is 2.29 excluding this outlier); (D) exons 2 and 4. Approximate fold change (≥ 2) is shown only for significant pairwise comparisons between NOD and other mouse strains ($P < 0.05$, adjusted for multiple testing). Bars, mean expression level (\pm pooled SEM for each tissue).

2/3 splice product, suggesting that this microdeletion may shorten the intron in which it resides, impeding the splicing lariat structure and altering expression of splice isoforms (Black 2003). However, increased splicing of these two exons was not sufficient to increase T1D protection (e.g., *Idd11E* mice develop T1D similar to NOD mice but do not harbor the NOD microdeletion and have increased expression of the exon 2/3 splice product). Further comparison of our congenic strains also indicated that only *Idd11D*, which has the greatest T1D protection, exhibited significantly increased expression of *AK005651* for all three spliced products in both the thymus and spleen compared with NOD. *Idd11G* and *Idd11F* have the next best T1D protection, respectively, but lack a similar increase and correlation between *AK005651* expression and T1D protection. This lack of correlation may reflect that sequence variation within the *Idd11* haplotype also affects the function of the encoded gene product (whether a large noncoding RNA or protein) leading to increased T1D protection, especially in combination with altered gene expression. The low level of expression for *AK005651*, although seemingly technically challenging (e.g., Ct values > 30 using the maximum amount of RNA/cDNA), was reproducibly detected by three different real-time PCR assays in different mouse cohorts. Given the relatively small differences observed between congenic strains, further studies are required to determine the function of *AK005651* and how sequence composition within the *Idd11* haplotype affects gene function and

expression to alter T1D susceptibility. The recent availability of efficient NOD ES cells (Nichols et al. 2009) should also enable replacement of the NOD variants with B6-derived protective variants to ultimately confirm the effect of this apparent gene-based functional haplotype. At present, *AK005651* represents a novel disease susceptibility gene compared with the current set of identified *Idd* genes, which mainly encode proteins of the immune system (e.g., MHC molecules, B2M, IL2, IL21, CTLA4, SLC11A1) that were relatively well characterized before their discovery as T1D genes (Serreze and Leiter 2001; Ridgway et al. 2008).

Discovery of *Idd11* relied on the presence of a recombination hotspot. The ability to map a disease locus using congenic mice depends on the occurrence of recombination events to generate successively smaller genomic intervals to identify the underlying gene. However, recombination during meiosis is not random (Petes 2001; Kauppi et al. 2004). Recombination hotspots and associated flanking coldspots ultimately restrict how small the congenic interval can become and dictate the number of candidate genes to be investigated. Paradoxically, *Idd11* mapped to a recombination hotspot. Although this hotspot demonstrated lower crossover activity compared with other hotspots within the mouse genome (Guillon and de Massy 2002; Yauk et al. 2003; Kauppi et al. 2007; Paigen et al. 2008), it was highly localized (~ 1.2 kb)

and resulted in one instance of a crossover with a complex conversion tract. This recombination hotspot provided the molecular mechanism that generated the “hybrid haplotypes” in our congenic NOD strains. It seems likely it is also responsible for the underlying allelic variation at this locus within the *Mus* species.

Rather than restricting our ability to map *Idd11*, this recombination hotspot enabled us to identify specific variants accounting for the varying levels of T1D susceptibility observed in our congenic strains. For example, the noncrossover/conversion event at rs27507073 (variant 3 in the *Idd11* haplotype) significantly increased the risk of T1D in *Idd11E* mice compared with *Idd11G* mice. Gene conversion, as well as genomic rearrangement, associated with recombination hotspots has been implicated in a number of Mendelian diseases (Lupski and Stankiewicz 2005; Chen et al. 2007; Turner et al. 2008). Our congenic strains demonstrate that recombination hotspots can generate unique hybrid haplotypes, due to shuffled haplotypes and/or complex conversion tracts, which increase the risk for a complex genetic disease if they arise on a genetic background of some liability. Up to 85% of T1D cases in the human population are sporadic (Karvonen et al. 2001). Undoubtedly these affected individuals inherited some combination of diabetogenic alleles from their parents, but common variants (including copy number variation) identified in recent genome-wide association studies have failed to explain the total T1D risk attributable to genetic factors (Concannon et al.

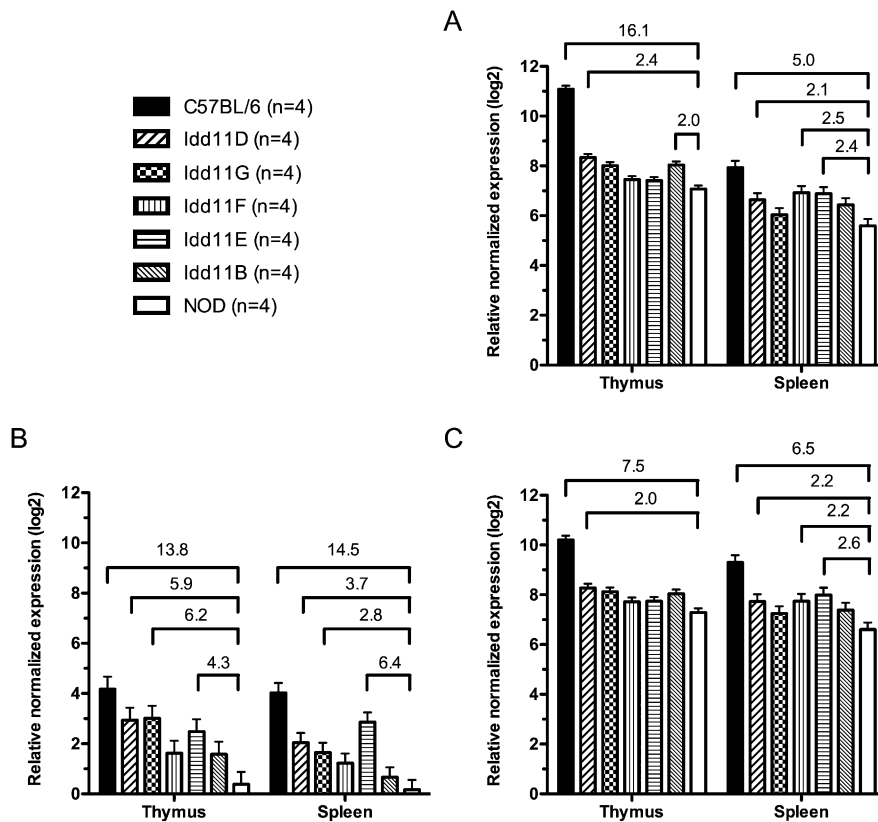


Figure 4. Expression analysis of *AK005651* in congenic NOD mouse strains. Quantitative real-time PCR was performed to detect expression differences between mouse strains at 50 d of age for the spliced product derived from the following: (A) exons 4 and 5, which are present in both splice variants; (B) exons 2 and 3; (C) exons 2 and 4. Different cohorts of NOD, B6, and Idd11D mice to those in Figure 3 were generated and used in conjunction with Idd11B, Idd11E, Idd11F, and Idd11G mice for this experiment. Fold change (≥ 2) is shown only for significant pairwise comparisons between NOD and other mouse strains ($P < 0.05$, adjusted for multiple testing). Bars, mean expression level (\pm pooled SEM for each tissue).

2009b; Conrad et al. 2009; The Wellcome Trust Case Control Consortium 2010). Instead, recombination hotspots may contribute to the generation of rare variants and/or hybrid haplotypes accounting for the significantly increased T1D risk in those individuals who have inherited a certain level of genetic liability due to common variants.

Our study represents a primary example of a recombination hotspot associated with a complex genetic disease. Ng et al. (2010) have also recently described a recombination hotspot in a region of *GABRB2* for which haplotypes are associated with schizophrenia. Similar to our study in congenic mice, their human study suggests that recombination hotspots are likely to contribute to the etiology of complex genetic diseases (Ng et al. 2010). However, sequence variation for the region on the human chr 1p35 encompassing the homologous *Idd11* locus has not been associated with T1D (Barrett et al. 2009). This is not necessarily unexpected as the NOD mouse represents a “single case study” with a collection of diabetogenic alleles, some of which are unique to the NOD mouse strain, while others are common within the *Mus* species (Atkinson and Leiter 1999; Serreze and Leiter 2001). Given that the NOD *Idd11* haplotype is unique, the equivalent susceptibility locus in humans might not exist or the equivalent human allele(s) may not be detectable in recent association studies because it is rare and/or resides in a recombination hotspot. HapMap data phase III (release 2, Febru-

ary 2009) shows that the homologous interval on the human chr 1p35 (between rs16834708 and rs13426) exhibits weak to no linkage disequilibrium ($r^2 < 0.1$) (The International HapMap Consortium 2007). Such potential recombination hotspots ($\sim 33,000$ estimated in the human genome) (The International HapMap Consortium 2007) have so far been neglected in genetic studies due to the statistical methods employed for detecting association (Kauppi et al. 2004; The International HapMap Consortium 2007), but their ability to give rise to gene conversion and unique hybrid haplotypes, as observed for the *Idd11* locus, demonstrates their capacity for producing sequence variation with demonstrable effects upon complex traits.

Methods

Mice

NOD/Lt (NOD) and C57BL/6 (B6) mouse strains were obtained from The Walter & Eliza Hall Institute specific pathogen-free (SPF) facilities. NOD.B6Idd11A, NOD.B6Idd11B, NOD.B6Idd11C, and NOD.B6Idd11D were established after 10 back-cross generations or more using a conventional breeding approach for congenic mouse strains as previously described (Rogner and Avner 2003; Brodnicki et al. 2005). NOD.B6Idd11E, NOD.B6Idd11F, and NOD.B6Idd11G mouse strains were generated from (NOD \times NOD.B6Idd11D) F_2 progeny (Supplemental Fig. 1), which were screened for recombination events between *D4Mit72* and *D4Mit204*. New congenic intervals, which dissected the Idd11D interval (Fig. 1), were fixed to homozygosity by brother–sister mating.

Genotyping and sequencing

DNA samples were extracted from tail biopsies by standard methods and genotyped with polymorphic markers by PCR (Supplemental Table 1; Brodnicki et al. 2000). NOD.B6Idd11D was genotyped using a 10-cM averaged genome-wide marker panel, and no B6-derived alleles were found outside the congenic interval (note that all other congenic strains described were derived from NOD.B6Idd11D). To fine-map the recombination sites, new genetic markers (i.e., nucleotide repeats) were identified using the publicly available mouse genome sequence (NCBI Build 37 assembly; mm9), the UCSC Genome Browser (<http://genome.ucsc.edu/>) (Kuhn et al. 2009), and the Tandem Repeats Finder program (Benson 1999). These markers (*D4Wehi1–D4Wehi22*) were shown to be polymorphic between the NOD and B6 mouse strains by PCR and gel electrophoresis of genomic DNA using sequence-specific oligonucleotides (Supplemental Table 1). Sequence within the defined *Idd11* critical interval was determined by direct sequencing of overlapping PCR products (Supplemental Table 2) and the BigDye Terminator v3.1 sequence kit (Applied Biosystems). Sequence contigs for each strain were aligned to determine sequence

variation between inbred strains (Megalyn, DNASTar, Inc.). Genotyping of sequence variants 1–8 within the *Idd11* haplotype (Table 2) was performed on a Roche LightCycler 480 using TaqMan probes (Applied Biosystems) (Supplemental Table 3) and Roche Probe Master Mix (Roche Applied Science) according to manufacturer's instructions.

Diabetes monitoring

Cohorts of female mice were housed in an SPF facility and tested once a week for elevated urinary glucose (>110 mmol/L) using Diastix reagent strips (Bayer Australia, Ltd.) over a 300-d time course. Three consecutive elevated readings indicated the onset of diabetes. Pairwise comparisons of the diabetes incidence between mouse strains were done using the log-rank test.

Bioinformatic analysis of *AK005651*

Genomic sequence and transcript sequences were aligned to the mouse genome (NCBI Build 37 assembly; mm9) using BLAT via the UCSC Genome Browser (<http://genome.ucsc.edu/>) (Kuhn et al. 2009). Sequence was compared with annotated ESTs, multispecies alignments, conserved elements, and the gene prediction algorithms available through the UCSC Genome Browser. Transcript sequences were also aligned to the NCBI EST database using BLASTN and to the NR protein database using BLASTX (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>; Altschul et al. 1990). All potential transcripts were translated in six frames and searched for known domains represented by profile hidden Markov models from the PFAM database using hmmpfam (<http://hmmer.janelia.org>) (Finn et al. 2008). The RFAM database was used to evaluate the presence of a noncoding gene (<http://rfam.janelia.org/>; Gardner et al. 2009), and BLASTN was used to align transcript sequence to mature miRNAs and stem-loop sequences in miRBase (Griffiths-Jones et al. 2006). To determine if variants are within predicted TFBSs, 71 nucleotides (nt) of sequence centered on the individual variants were extracted from the mouse genome, and TFBSs were predicted in these sequences using only high-quality TRANSFAC matrices (Matys et al. 2006) representing vertebrate transcription factors and match score thresholds selected to give the minimum false-positive rate (Kel et al. 2003).

Quantitative real-time PCR

Tissues were taken from female mice (~50 d old) and RNA isolated using TRIzol reagent (Invitrogen). cDNA was synthesized using Superscript III reverse transcriptase (Invitrogen). Quantitative real-time PCR was performed on a Roche LightCycler 480 using LightCycler Probe Master Reagent (Roche Diagnostics) or TaqMan Gene Expression Master Mix (Applied Biosystems) according to the manufacturer's instructions. Technical replicates were done in triplicate to calculate the average Ct value for each biological sample. Oligonucleotide primers and fluorescent probes were synthesized by Sigma Genosys, Roche Applied Sciences, or Applied Biosystems (Supplemental Table 9). The fluorescent probes used to detect the spliced products between *AK005651* exon 2 and exon 3, as well as exon 2 and exon 4, are located across the spliced exon boundaries, whereas the fluorescent probe used to detect the spliced product between exon 4 and exon 5 is located entirely within exon 5 (Fig. 3) or across the spliced exon boundary (Fig. 4). The fluorescent probe used to detect the spliced product for *Spocd1* was located in exon 9. The fluorescent probe used to detect the spliced product for *Bai2* was located in exon 29. Thermal cycling consisted of a denaturation step (10 min at 95°C) and 45 amplification cycles (10 sec at 95°C, 15–30 sec at 60°C–66°C, 30 sec at

40°C). Products observed for each primer pair were confirmed by sequencing. Standard curves were generated for all primer sets to ensure exponential increase of targeted transcripts during amplification (efficiency = $10^{(-1/\text{slope})} = \sim 2$). ΔCt for each tissue was calculated as $\text{Ct}^{\text{AK005651}} - \text{Ref}$, where Ref is the average Ct value of the reference genes *Hmbs* and *Hprt1*. Relative normalized log₂ expression values were calculated for graphing purposes as $41 - \text{mean}(\text{Ref}) - \Delta\text{Ct}$, where mean(Ref) is the grand mean of the reference genes Ct for each tissue. Here, 40 represents the practical Ct detection limit for real-time PCR, and 41 establishes the 0 point (i.e., no detectable gene expression) for the y-axis. Subtracting mean(Ref) converts the scores to log₂-expression relative to the detection threshold scale from 0–41. Statistical significance for the difference in expression was obtained using pairwise *t*-tests with pooled standard deviations for each tissue. *P*-values were adjusted for multiple testing using Holm's method (Holm 1979).

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