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### Published In/Presented At

Reed, D. R., Li, S., Li, X., Huang, L., Tordoff, M. G., Starling-Roney, R., & ... Bachmanov, A. A. (2004). Polymorphisms in the taste receptor gene (Tas1r3) region are associated with saccharin preference in 30 mouse strains. *The Journal Of Neuroscience: The Official Journal Of The Society For Neuroscience*, 24(4), 938-946.

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# Polymorphisms in the Taste Receptor Gene (*Tas1r3*) Region Are Associated with Saccharin Preference in 30 Mouse Strains

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The results of recent studies suggest that the mouse *Sac* (saccharin preference) locus is identical to the *Tas1r3* (taste receptor) gene. The goal of this study was to identify *Tas1r3* sequence variants associated with saccharin preference in a large number of inbred mouse strains. Initially, we sequenced ~6.7 kb of the *Tas1r3* gene and its flanking regions from six inbred mouse strains with high and low saccharin preference, including the strains in which the *Sac* alleles were described originally (C57BL/6J, *Sac<sup>b</sup>*; DBA/2J, *Sac<sup>d</sup>*). Of the 89 sequence variants detected among these six strains, eight polymorphic sites were significantly associated with preferences for 1.6 mM saccharin. Next, each of these eight variant sites were genotyped in 24 additional mouse strains. Analysis of the genotype–phenotype associations in all 30 strains showed the strongest association with saccharin preference at three sites: nucleotide (nt) –791 (3 bp insertion/deletion), nt +135 (Ser45Ser), and nt +179 (Ile60Thr). We measured *Tas1r3* gene expression, transcript size, and T1R3 immunoreactivity in the taste tissue of two inbred mouse strains with different *Tas1r3* haplotypes and saccharin preferences. The results of these experiments suggest that the polymorphisms associated with saccharin preference do not act by blocking gene expression, changing alternative splicing, or interfering with protein translation in taste tissue. The amino acid substitution (Ile60Thr) may influence the ability of the protein to form dimers or bind sweeteners. Here, we present data for future studies directed to experimentally confirm the function of these polymorphisms and highlight some of the difficulties of identifying specific DNA sequence variants that underlie quantitative trait loci.

**Key words:** sweet; preference; phenotype; genotype; haplotype; QTL

## Introduction

Mouse strains differ in their avidity to consume sweet solutions (Pelz et al., 1973; Stockton and Whitney, 1974; Ramirez and Fuller, 1976; Lush, 1989). Fuller (1974) demonstrated that most of the differences in saccharin preferences between C57BL/6J and DBA/2J strains depend on a single locus, *Sac*, with the dominant *Sac<sup>b</sup>* allele present in the C57BL/6J strain. Subsequent studies confirmed this finding in the BXD recombinant inbred strains, in crosses between the C57BL/6 and DBA/2 strains (Lush, 1989; Belknap et al., 1992; Phillips et al., 1994; Lush et al., 1995; Blizard et al., 1999) and in crosses between the C57BL/6ByJ and 129P3/J strains (Bachmanov et al., 1997). The *Sac* locus has been mapped to the subtelomeric region of mouse chromosome 4 (Phillips et al., 1994; Lush et al., 1995; Bachmanov et al., 1997; Blizard et al., 1999; Li et al., 2001) and more recently has been cloned position-

ally (Bachmanov et al., 2001). It corresponds to the *Tas1r3* gene, which encodes a sweet taste receptor, T1R3 (Kitagawa et al., 2001; Max et al., 2001; Montmayeur et al., 2001; Nelson et al., 2001; Sainz et al., 2001; Li et al., 2002a; Nelson et al., 2002; Ariyasu et al., 2003; Damak et al., 2003).

Several studies have attempted to identify polymorphisms within the *Tas1r3* gene that are equivalent to allelic variants of the *Sac* locus, by comparing *Tas1r3* sequences in mouse strains with the most extreme differences in sweetener preferences (Bachmanov et al., 2001; Kitagawa et al., 2001; Max et al., 2001; Montmayeur et al., 2001; Sainz et al., 2001). However, saccharin intake in mice is a quantitative trait, and the *Sac* locus accounts for a different proportion of behavioral variation in different mouse crosses (Lush et al., 1995; Bachmanov et al., 1997; Blizard et al., 1999). The current study was designed to examine saccharin preference as a quantitative trait. First, we compared many inbred mouse strains with distant genealogies. Second, we examined genomic sequences including *Tas1r3* exons, introns, and upstream and downstream regions, so that polymorphisms affecting amino acid composition, RNA splicing, or potential regulatory regions could be detected. Finally, we statistically compared the strength of the associations between the phenotype and genotype. The influence of these polymorphisms on *Tas1r3* gene ex-

Received May 7, 2003; revised Nov. 3, 2003; accepted Nov. 19, 2003.

This work was supported by National Institutes of Health Grants R03DC03509, R01DC04188, and R01DK55853 (D.R.R.); R03DC05154 (L.H.); R01AA11028 (M.G.T.); R01DC00882 (G.K.B.); and R03DC03854 (A.A.B.). Maria Theodorides, Lynn Vo, and Kirsten Mascioli provided technical assistance.

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DOI:10.1523/JNEUROSCI.1374-03.2004

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pression in taste tissue was evaluated by comparing two mouse strains with different *Tas1r3* haplotypes and phenotypical *Sac* alleles.

## Materials and Methods

**General approach.** The analysis of the associations of *Tas1r3* sequence variants with saccharin preference was conducted in two stages. First, we selected three strains with strong saccharin preferences (C57BL/6J, CAST/Ei, and SWR/J) and three strains indifferent to saccharin (AKR/J, DBA/2J, and 129P3/J). In these six strains, an ~6.7 kb segment of genomic DNA, including ~2.4 kb upstream and ~1.0 kb downstream of *Tas1r3*, was sequenced. The analysis of saccharin preferences and sequence variants in these strains identified eight significant associations. For the second stage, genomic regions encompassing these eight variants were examined in an additional 24 strains of mice, and genotype–phenotype associations were then analyzed in all 30 strains.

**Sweet preference tests.** Mice from 28 inbred strains (129P3/J, A/J, AKR/J, BALB/cByJ, BUB/BnJ, C3H/HeJ, C57BL/6J, C57L/J, CAST/Ei, CBA/J, CE/J, DBA/2J, FVB/NJ, I/LnJ, KK/HIJ, LP/J, NOD/LtJ, NZB/BINJ, P/J, PL/J, RBF/DnJ, RF/J, RIIS/J, SEA/GnJ, SJL/J, SM/J, SPRET/Ei, and SWR/J) were purchased from The Jackson Laboratory (Bar Harbor, ME) and used in two-bottle preference tests with 1.6 mM saccharin and water. Because mice from the IS/Cam and ST/bJ strains used for *Tas1r3* sequencing were not available from The Jackson Laboratory, we used 1.6 mM saccharin preference data for these strains obtained from a previous experiment that used a similar method (Lush, 1989). To justify combining the data from two different experiments, we analyzed correlations between 1.6 mM saccharin preference scores for a subset of 15 strains used in both studies. The correlation coefficient for strain means obtained in our study and in the study by Lush was  $r = +0.62$  ( $p = 0.013$ ). Considering that different substrains were used in these two studies, and that genetic drift within strains could occur during the  $\geq 11$  years separating these two experiments, this result demonstrates consistency between laboratories.

The construction of drinking tubes and the testing procedure were described in detail previously (Bachmanov et al., 1996b). Groups of individually caged, 2- to 3-month-old male mice [ $n = 12$  for each strain, except CAST/Ei, BUB/BnJ, NZB/BINJ, and RBF/DnJ ( $n = 11$ ) and BALB/cByJ ( $n = 10$ )] were offered for 48 hr two drinking tubes, one containing 1.6 mM saccharin and one containing water. The tube positions were reversed after 24 hr. For each 24 hr period, the volumes of water and solution consumed were recorded. Preference scores were calculated for each mouse as the average daily amount of saccharin consumed divided by the average daily total amount of fluid (saccharin plus water) consumed, multiplied by 100.

**DNA sequencing and genotyping.** Genomic DNA from 31 inbred mouse strains (129P3/J, A/J, AKR/J, BALB/cByJ, BUB/BnJ, C3H/HeJ, C57BL/6ByJ, C57BL/6J, C57L/J, CAST/Ei, CBA/J, CE/J, DBA/2J, FVB/NJ, I/LnJ, IS/CamEi, KK/HIJ, LP/J, NOD/LtJ, NZB/BINJ, P/J, PL/J, RBF/DnJ, RF/J, RIIS/J, SEA/GnJ, SJL/J, SM/J, SPRET/Ei, ST/bJ, and SWR/J) was purchased from the Mouse DNA Resource at The Jackson Laboratory. Sequences of the *Tas1r3* region of the C57BL/6ByJ strain [used in our genetic mapping studies to clone positionally the *Sac* locus (Bachmanov et al., 1997, 2001; Li et al., 2001)] and the C57BL/6J strain (used in the strain survey of saccharin preferences described above) were identical, and so this sequence is referred to as C57BL/6 in this study.

Sixteen primer pairs were designed to encompass the 6675 bp *Tas1r3* region, including 2482 bp upstream of the putative methionine initiation codon and 1046 bp downstream of the putative termination codon (primer sequences and PCR conditions are available on request). These upstream and downstream regions include most of the intergenic intervals between the *Tas1r3* coding region and two flanking genes, *Dvl* and *BC002216* (GenBank accession number NM\_024472) (Bachmanov et al., 2001; Li et al., 2002b). Genomic DNA was used as a template to generate PCR products, which were sequenced. The majority of sequencing was conducted at Pfizer Global Research and Development (Alameda, CA) with additional sequencing conducted by the University of Pennsylvania Sequencing Facility (Philadelphia, PA). The Sequencher software pro-

gram (Genecodes, Ann Arbor, MI) was used to assemble and align sequences (GenBank accession numbers AF550928 to AF551332) and to identify sequence variants. To define nucleotide (nt) positions of the variable sites, we aligned the C57BL/6 genomic sequences of *Tas1r3* (GenBank accession number AF311386) and BAC RPC123-118E21 (GenBank accession number AF389853). Polymorphism sites were numbered two ways. First, polymorphism positions were numbered using the putative start codon of *Tas1r3* as +1. Second, because of the large number of insertions and deletions among inbred strains, polymorphism positions were also given relative to their position in BAC 118E21. To identify the correct nucleotides at position +661 in the A/J strain and at position +1309 in the 129/P3J strain, we consulted the Celera Discovery System. The deletion/insertion variable sites at nt -791 and +3479 were genotyped using simple sequence length polymorphism and single-stranded conformational polymorphism methods.

**Proportion of transitions and transversions.** The ratio of transitions and transversions found in the *Tas1r3* gene was compared with the rates of nucleotide substitutions in the whole mouse genome using a  $\chi^2$  test [ $>39,000$  single nucleotide polymorphisms (SNPs); Mouse Phenome Database].

**Statistical analyses of genotype–phenotype relationships.** Preference scores of individual mice were averaged to obtain means for each strain, and the means were used in the statistical analyses. The relationship between *Tas1r3* sequence variants and saccharin preferences was analyzed using an approach similar to a previous study (Long et al., 1998). The first stage of the analysis was conducted using all polymorphisms detected in six strains (129P3/J, AKR/J, C57BL/6J, CAST/Ei, DBA/2J, and SWR/J). For each polymorphic site, the effect of the nucleotide variant on saccharin preference was estimated using one-way ANOVA. For this initial stage of analysis, we have chosen arbitrarily the threshold of statistical significance to be  $p < 0.01$  as a compromise between the standard threshold of  $p < 0.05$  (which could produce approximately five false positives for 89 comparisons made here) and a Bonferroni correction ( $p < 0.05/89 \approx 0.00056$ ), which would be too conservative in the initial stage of analysis using a small number of strains. All polymorphisms analyzed in the second stage had only two variants among the 30 strains; therefore, we used *t* tests to analyze genotype–phenotype relationships. The strains were grouped by sequence variant at each of the eight polymorphic sites, and the saccharin preferences of the two groups of strains were compared by two-tailed *t* test. The obtained *p* values were changed to their negative base-10 logarithms for ease of presentation, where needed. All analyses were done using Statistica software (StatSoft, Tulsa, OK).

Heritability of saccharin preference was calculated as the ratio  $SS_{\text{among strains}}/SS_{\text{total}}$  based on the sums of squares (SS) obtained in a one-way ANOVA assessing the effect of strain and using preference scores of individual mice (Belknap, 1998). Similarly, the percentage of genetic variation explained by alleles at the *Tas1r3* polymorphic sites was calculated as the ratio  $SS_{\text{Tas1r3 genotype}}/SS_{\text{total}}$  based on the results of a one-way ANOVA assessing the effect of the *Tas1r3* genotype and using the mean preference scores for the mouse strains.

**Bioinformatic analyses of DNA variants.** We used several approaches to identify the potential biological significance of the eight variant sites associated with saccharin preferences. The three variant sites residing within coding regions were examined to determine whether they were synonymous (encoded the same amino acid) or nonsynonymous (encoded different amino acids). The variant sites upstream of the putative methionine initiation codon and the variant site downstream of the putative termination codon were examined using several methods. First, the sequences were analyzed to determine whether the variants were contained within repetitive elements, using the computer program Repeatmasker (A. F. A. Smit and P. Green, unpublished results). Second, each variant site and the flanking region was examined for alterations in putative transcription factor-binding sites. We used the computer program MatInspector 2.2 (Quandt et al., 1995) in conjunction with the Transfac 4.0 database of transcription factors and their binding sites (Wingender et al., 2000). Third, the sequenced upstream and downstream regions of all three members of the mouse *Tas1r* family [*Tas1r1* (GenBank accession number AL611927), *Tas1r2*, and *Tas1r3* (GenBank accession number AF389853)] were aligned, and conserved elements

**Table 1. Polymorphisms within the *Tas1r3* region of mouse strains with high (C57BL/6, CAST/Ei, and SWR/J) and low (129P3/J, AKR/J, and DBA/2J) saccharin preferences**

Position <sup>a</sup>	AF389853 <sup>b</sup>	Location	C57BL/6 <sup>c</sup>	SWR/J	CAST/Ei	129P3/J	DBA/J	AKR/J
–2406	50853	5'	G	T	T	T	T	T
<b>–2386</b>	<b>50832</b>	<b>5'</b>	<b>T</b>	<b>T</b>	<b>T</b>	<b>A</b>	<b>A</b>	<b>A</b>
–2343	50790	5'	C	DEL	C	DEL	DEL	DEL
–2342	50789	5'	A	G	G	G	G	G
–2330	50777	5'	A	G	A	G	G	G
–2323	50770	5'	A	G	G	G	G	G
–2315	50762	5'	G	DEL	G	DEL	DEL	DEL
–2198	50654	5'	DEL	T	T	T	T	T
–2181	50629	5'	G	T	T	T	T	T
–2002	50450	5'	C	T	T	T	T	T
–1994	50442	5'	C	A	A	A	A	A
–1845	50293	5'	A	G	A	G	G	G
–1820	50268	5'	G	A	G	A	A	A
–1803	50251	5'	G	A	G	A	A	A
–1791	50239	5'	C	T	C	T	T	T
–1786	50234	5'	G	A	A	A	A	A
–1558	50006	5'	G	UN	A	UN	G	G
–1488	49936	5'	A	UN	G	UN	G	G
–1417	49865	5'	A	UN	G	UN	G	G
–1414	49863	5'	DEL	UN	TT	UN	TT	TT
–1407	49856	5'	C	UN	T	UN	T	T
–1317	49767	5'	G	UN	A	G	G	G
–1284	49734	5'	G	UN	G	G	A	A
–1273	49723	5'	A	A	G	A	A	A
–1260	49710	5'	C	C	A	C	C	C
–1200	49650	5'	A	T	A	T	T	T
–1130	49580	5'	A	G	A	G	G	G
–1078	49529	5'	A	G	G	G	G	G
–1032	49482	5'	T	G	G	G	G	G
–967	49417	5'	C	T	T	T	T	T
–962	49412	5'	T	C	C	C	C	C
–912	49362	5'	C	C	T	C	C	C
–827	49277	5'	AG	AG	DEL	AG	AG	AG
–822	49272	5'	A	A	G	A	A	A
–817	49267	5'	A	G	G	G	G	G
<b>–791</b>	<b>49241</b>	<b>5'</b>	<b>AGG</b>	<b>AGG</b>	<b>AGG</b>	<b>DEL</b>	<b>DEL</b>	<b>DEL</b>
–753	49203	5'	T	T	C	T	T	T
–564	49014	5'	G	C	G	C	C	C
–349	48799	5'	T	C	T	T	T	T
–320	48770	5'	12 bp	DEL	12 bp	DEL	DEL	DEL
–267	48718	5'	G	DEL	A	DEL	DEL	DEL
–162	48613	5'	G	C	C	C	C	C
<b>–138</b>	<b>48588</b>	<b>5'</b>	<b>A</b>	<b>A</b>	<b>A</b>	<b>G</b>	<b>G</b>	<b>G</b>
–104	48554	5'	G	G	A	G	G	G
–32	48482	5'	C	T	C	T	T	T
–30	48480	5'	C	T	T	T	T	T
–9	48459	5'	C	A	C	C	C	C
<b>135</b>	<b>48316</b>	<b>Exon 1</b>	<b>A</b>	<b>A</b>	<b>A</b>	<b>G</b>	<b>G</b>	<b>G</b>
<b>163</b>	<b>48288</b>	<b>Exon 1</b>	<b>A</b>	<b>A</b>	<b>A</b>	<b>G</b>	<b>G</b>	<b>G</b>
<b>179</b>	<b>48272</b>	<b>Exon 1</b>	<b>T</b>	<b>T</b>	<b>T</b>	<b>C</b>	<b>C</b>	<b>C</b>
182	48269	Exon 1	C	T	C	T	T	T
186	48265	Exon 1	C	T	T	T	T	T
264	48187	Intron	DEL	G	G	G	G	G
271	48180	Intron	A	G	G	G	G	G
272	48179	Intron	G	G	A	G	G	G
319	48139	Exon 2	T	C	T	C	C	C
556	47902	Exon 2	A	A	G	A	A	A
<b>661</b>	<b>47799</b>	<b>Intron</b>	<b>T</b>	<b>T</b>	<b>T</b>	<b>C</b>	<b>C</b>	<b>C</b>
701	47759	Exon 3	T	C	C	C	C	C
974	47486	Exon 3	A	G	G	G	G	G
978	47482	Exon 3	C	T	C	C	C	C
1309	47151	Exon 3	G	A	A	A	A	A
1466	46994	Exon 3	C	T	C	C	C	C

(Table continues)

**Table 1. Continued**

Position <sup>a</sup>	AF389853 <sup>b</sup>	Location	C57BL/6 <sup>c</sup>	SWR/J	CAST/Ei	129P3/J	DBA/J	AKR/J
1874	46586	Intron	A	A	DEL	A	A	A
1897	46563	Intron	A	C	C	C	C	C
2061	46399	Intron	G	G	DEL	G	G	G
2098	46362	Intron	G	G	G	C	G	G
2106	46354	Intron	A	A	G	A	A	A
2216	46244	Exon 6	A	A	G	A	A	A
2405	46055	Exon 6	T	C	C	C	C	C
2474	45985	Exon 6	G	A	A	A	A	A
2531	45929	Exon 6	A	A	G	A	A	A
2651	45809	Exon 6	A	A	T	A	A	A
2656	45762	Exon 6	T	C	T	T	T	T
2698	45762	Exon 6	T	A	T	C	C	C
3145	45315	Exon 6	G	A	A	A	A	A
3240	45220	3'	G	G	A	G	G	G
3434	45026	3'	T	C	T	C	C	C
<b>3479</b>	<b>44981</b>	<b>3'</b>	<b>A</b>	<b>A</b>	<b>A</b>	<b>DEL</b>	<b>DEL</b>	<b>DEL</b>
3483	44977	3'	C	T	C	T	T	T
3493	44967	3'	C	G	G	G	G	G
3581	44870	3'	(TG) <sub>23</sub>	(TG) <sub>15</sub>	(TG) <sub>20</sub>	(TG) <sub>23</sub>	(TG) <sub>16</sub>	(TG) <sub>16</sub>
3609	44851	3'	G	A	G	G	G	G
3693	44767	3'	T	T	C	T	T	T
3724	44736	3'	T	T	C	T	T	T
3737	44723	3'	G	A	G	G	G	G
3781	44679	3'	C	T	C	T	T	T
3782	44678	3'	A	G	A	G	G	G
4201	44259	3'	A	G	A	G	G	G

The 12 bp sequence at polymorphic position –320 is GCAGCTTCITTT. The dinucleotide repeat at position +3581 is imperfect in the C57BL/6 and SWR/J strains, with G substituted with A in the second TG repeat.

DEL, Deletion; UN, unknown. Polymorphisms significantly associated with saccharin preference are shown in bold.

<sup>a</sup>Polymorphism positions relative to the first nucleotide in the putative start codon of *Tas1r3*.

<sup>b</sup>Position in genomic sequence (GenBank accession number AF389853).

<sup>c</sup>Sequences of the C57BL/6J and C57BL/6ByJ strains were identical and shown here as C57BL/6.

were identified based on the sequence similarity assessed using a nucleic acid dot plot (Maizel and Lenk, 1981). Similarly, to identify elements conserved between human *TAS1R3* (GenBank accession number AL139287) and mouse *Tas1r3* (GenBank accession number AF389853) genes, the upstream and downstream regions of these genes were aligned using PipMaker (Schwartz et al., 2000). Then we determined whether these regions conserved among mouse *Tas1r* genes or between mouse and human *Tas1r3* orthologs contained any of the eight sequence variants associated with saccharin preference. By examining the exon/intron boundaries, we determined whether any variant site interrupted a splice junction site. Finally, gene promoter regions were predicted using the computer program Promoter 2.0 (Knudsen, 1999), and we determined whether any of the eight DNA sequence variants were contained within these sites.

**Subjects and tissues used to assess *Tas1r3* gene expression.** We tested the hypothesis that the polymorphisms that predict saccharin intake act by altering *Tas1r3* gene expression, transcript size, or protein abundance in taste tissue. Lingual tissues used for real-time PCR, Northern blot hybridization, and immunocytochemistry were collected from killed C57BL/6J (B6) and 129P3/J (129) mice. Tissues containing fungiform, foliate, and circumvallate taste papillae and control tissue (lingual epithelium without taste papillae) were dissected as described previously (Huang et al., 1999).

**Northern blot hybridization.** Isolation of mouse total RNA from circumvallate and foliate taste papillae, electrophoresis, transfer, fixation, hybridization, washing, and exposure were as described (Huang et al., 1999), with slight modifications. The <sup>32</sup>P-labeled *Tas1r3* probe was generated by random nonamer priming of a 1.3 kb cDNA fragment of mouse T1R3 corresponding to the 5' end coding sequence using Exo (-) Klenow polymerase in the presence of (α-<sup>32</sup>P)-dCTP. Ethidium bromide-stained 18S and 28S ribosomal RNA served as a control for RNA quality and gel loading.

**Real-time PCR.** Total RNA was extracted from the lingual tissues of 10

mice (5 mice from each strain) using the Absolutely RNA Microprep kit (Stratagene, La Jolla, CA). First-strand synthesis of cDNA was conducted following the directions of the manufacturer (SuperScript III; Invitrogen, Carlsbad, CA). Transcribed cDNA (1  $\mu$ l/well) was transferred to a PCR 96-well optical reaction plate. Each well was supplemented with predeveloped TaqMan assay reagents (primer and fluorescent probe) and Universal PCR Master Mix to a final volume of 20  $\mu$ l. The PCR product was heated to 50°C for 2 min and 95°C for 10 min, and then 40 amplification cycles were conducted at 95°C for 15 sec and 60°C for 60 sec. DNA amplification was performed in an ABI PRISM 7000 Sequence Detection System (Applied Biosystems, Foster City, CA), detecting fluorescence (500–560 nm) after laser light excitation. PCR cycle time values represent the time at which the emitted fluorescence increased above threshold. Primers were designed to span an intron to reduce genomic DNA contamination. Genomic contamination was assessed by including additional samples in the assay that contained no reverse transcriptase. *Tas1r3* gene expression was normalized as described previously (Zehnder et al., 2003) to a ubiquitously expressed control gene (TATA box-binding protein; *Tbp*; GenBank accession number NM\_013684) and a taste receptor cell-specific gene involved in signal transduction (guanine nucleotide-binding protein 13, gamma; *Gng13*; GenBank accession number NM\_022422). Samples in which there was little or no housekeeping gene expression were excluded from additional analysis. The significance of differences in *Tas1r3* expression between inbred strains was assessed by a two-way ANOVA using tissue and strain as factors.

**Immunocytochemistry.** Polyclonal antisera against a hemocyanin-conjugated T1R3 peptide (aa 45–62) were raised in rabbits. Frozen sections of mouse lingual tissue (previously fixed in 4% paraformaldehyde and cryoprotected in 20% sucrose) were blocked in 3% BSA, 0.3% Triton X-100, 2% goat serum, and 0.1% sodium azide in PBS for 1 hr at room temperature and then incubated for 16 hr at 4°C with antisera against T1R3 (1:800). The secondary antibodies were Cy3-conjugated goat anti-rabbit Ig for T1R3. T1R3 immunoreactivity was blocked by preincubation of the antisera with the corresponding synthetic peptide at 10  $\mu$ M. Preimmune serum did not show any immunoreactivity.

## Results

### Saccharin preference

Responses to 1.6 mM saccharin in the two-bottle choice tests ranged from indifference (i.e., mice consumed similar amounts of saccharin and water and their preference scores were close to 50%) to a strong preference (i.e., mice consumed almost 100% of their total fluid intake as saccharin). The strain differences were highly significant ( $F_{(27,299)} = 25.7$ ;  $p < 10^{-16}$ ; one-way ANOVA) and heritable ( $h^2 = 0.70$ ). For the initial analysis of the genotype–phenotype associations, we selected three strains with strong saccharin preferences (C57BL/6, CAST/Ei, and SWR/J; preference scores,  $91 \pm 1$ ,  $97 \pm 1$ , and  $93 \pm 1\%$ , respectively) and three strains indifferent to saccharin (129P3/J, AKR/J, and DBA/2J; preference scores,  $58 \pm 2$ ,  $41 \pm 8$ , and  $52 \pm 2\%$ , respectively).

### Sequences of the *Tas1r3* region in six strains with high or low saccharin preferences

Comparison of DNA sequences of the *Tas1r3* region in the six inbred mouse strains with high or low saccharin preferences identified 89 sequence variants (Table 1), of which 86 (97%) had two alleles and 2 (2%) had three alleles. One polymorphism was an imperfect dinucleotide repeat with five alleles. Of all polymorphic sites, 77 contained SNPs (87% of polymorphic sites) and 12 contained deletions/insertions (13% of polymorphic sites); one polymorphic site with three alleles comprised both an SNP and a deletion/insertion. The DNA sequence variants in 58 cases (65%) occurred in one of six strains; in 21 cases (24%), they occurred in two of six strains, and in 9 cases (10%) they occurred in three of six strains. The proportions of transitions and transversions were similar to those of the whole mouse genome ( $\chi^2 = 0.32$ ; observed

**Table 2. Sequence variants within exons of the *Tas1r3* gene of six mouse strains**

Position (nt)	Codon	Amino acid
+135	TCA>TCG	Ser45Ser
+163	ACA>GCA	Thr55Ala
+179	ATC>ACC	Ile60Thr
+182	CCG>CTG	Pro61Leu
+186	TGC>TGT	Cys62Cys
+319	CTT>CTC	Leu68Leu
+556	TCA>TCG	Ser147Ser
+701	AGT>AGC	Ser168Ser
+974	GTA>GTG	Val259Val
+978	CGC>TGC	Arg261Cys
+1309	CGG>CAG	Arg371Gln
+1466	CAC>CAT	His423His
+2216	CCA>CCG	Pro545Pro
+2405	GGT>GGC	Gly608Gly
+2474	GGG>GGA	Gly616Gly
+2531	ACA>ACG	Thr650Thr
+2651	GTA>GTT	Val690Val
+2656	TTG>TCG	Leu692Ser
+2698	ATC>AAC>ACC	Ile706Asn706Thr
+3145	GGA>GAA	Gly855Glu

All nucleotide positions are relative to the putative ATG start codon of the *Tas1r3* gene. See Table 1 for sequence variants by inbred strain.

**Table 3. Frequencies (%) of *Tas1r3* polymorphisms among six mouse strains**

Strain	C57BL/6	CAST/Ei	SWR/J	129P3/J	AKR/J
CAST/Ei	57				
SWR/J	68	57			
129P3/J	<b>66</b>	<b>59</b>	<b>24</b>		
AKR/J	<b>68</b>	<b>57</b>	<b>23</b>	7	
DBA/2J	<b>68</b>	<b>56</b>	<b>20</b>	5	2

The percentage of variable sequence sites that were polymorphic within each pair of strains is shown. Pairings between high-preference and low-preference strains are shown in bold.

rates for each substitution: A or G = 34.6%, C or T = 34.3%, A or C = 8.5%, G or T = 8.5%, A or T = 8.2%, C or G = 6.0%). Of the nucleotide substitutions in protein coding regions, 8 of 20 (40%) were nonsynonymous (Table 2).

Frequencies of polymorphisms among strain pairs are shown in Table 3. There were substantial sequence differences between the high-prefering and low-prefering strains (24–68% of the 89 polymorphic sites). Sequence variability within the group of high-saccharin-prefering strains was higher (57–68%) than among the low-prefering strains (2–7%). The sequences of the low-prefering strains (129P3/J, AKR/J, and DBA/2J) were similar, although the strains are only related distantly (Beck et al., 2000). These three strains may have inherited this chromosomal region from a common ancestor, and the small percentage of polymorphisms among them may represent mutations accumulated during ~100 years of independent breeding of these strains.

### Polymorphisms associated with saccharin preferences in six strains

Of the 89 polymorphic sites, 8 sites were significantly associated with saccharin preference ( $F_{(1,4)} = 67.3$ ;  $p = 0.0012$ ) (Fig. 1). Each of the eight polymorphisms had only two sequence variants, one of which was found in all three high-saccharin-prefering strains, and the other one was found in all three low-prefering strains (Table 1). Therefore, these eight variable sites formed two haplotypes found either only in the high-saccharin-prefering strains or only in the low-saccharin-prefering strains.

Three of the eight polymorphisms significantly associated with saccharin preference were in the 5' region of *Tas1r3*, three

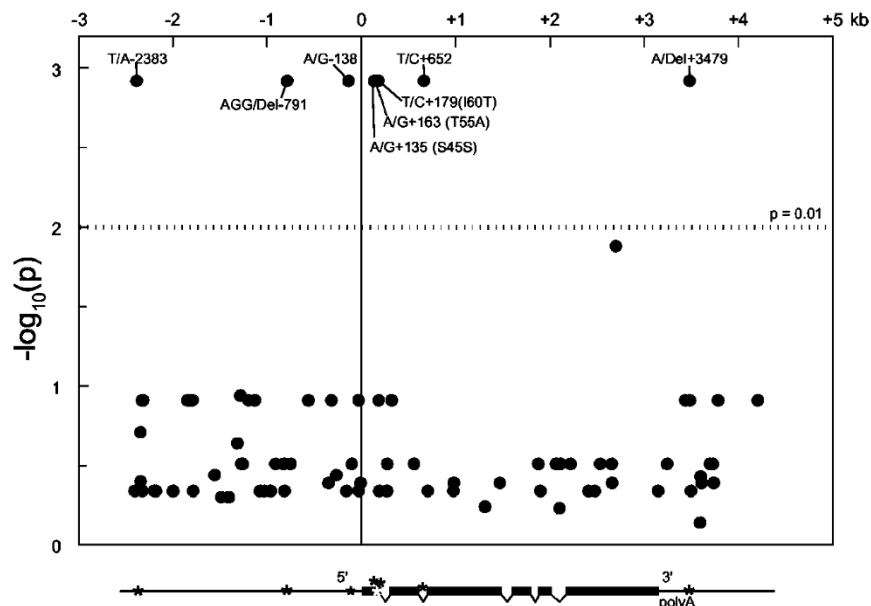
were in exon 1, one was in intron 2, and one was in the 3' region (Table 1). Among the three SNPs in exon 1, one was synonymous and two predicted an amino acid change.

Of the four variant sites located in the upstream and downstream regions, two sites did not alter putative transcription factor-binding sites. The SNP at nt -2383 in the high-preferring strains was a part of a predicted binding site for the transcription factor DPCR3HD, a cut-like homeodomain protein. In the low-preferring strains, this SNP was a part of a binding site for hepatocyte nuclear factor-3. The polymorphic site at nt -791 in strains with high-saccharin preferences was a part of a gut-enriched Krueppel-like factor-binding site, but a deletion at this position in the low-preferring strains eliminated this transcription factor-binding site. In addition, the -791 polymorphism was located at the beginning of a repetitive DNA element (long terminal repeat/mammalian retrotransposon like), but none of the other three variants were within repetitive elements.

The upstream and downstream DNA sequences had regions conserved between human *TAS1R3* and mouse *Tas1r3*, and there were also few stretches of conserved DNA sequence among the three members of the *Tas1r* family. However, none of the eight variant sites were found within these conserved regions. None of the eight DNA sequence variants interrupted an intron/exon boundary or a predicted promoter motif.

**Polymorphisms associated with saccharin preferences in 30 strains**

The genomic regions encompassing the eight polymorphisms significantly associated with saccharin preference in the six high- or low-saccharin-preferring strains were examined in an additional 24 inbred mouse strains, and genotype-phenotype associations were analyzed using all 30 strains. Twenty-six of the 30 strains had alleles of the eight polymorphic sites retained as the two haplotypes found in the high- and low-preferring strains (Table 4). Four strains (CE/J, I/LnJ, RF/J, and SM/J) had a broken haplotype, with three new combinations of genotypes at the polymorphic sites. Therefore, there were five unique strain distribution patterns of genotypes for the eight polymorphisms. As a result, analysis of the genotype-phenotype relationships allowed us to distinguish the polymorphic sites based on the strength of their associations with saccharin preferences (Fig. 2). All sites showed a highly significant association with saccharin preference ( $0.000065 \geq p \geq 0.00000000012$ ), with three variant sites (nt -791, nt +135, and nt +179) showing the strongest association. When inbred mouse strains were grouped by genotype at these three variant sites, there was a large difference in saccharin preference between groups. However, within each group, there was also substantial variation in preference (Fig. 3). Overall, variation in the *Tas1r3* haplotype for these three polymorphic sites explained 78% of the genetic variation in saccharin preferences among the inbred mouse strains; the remaining 22% of genetic variance is attributed to the contribution of other genes.



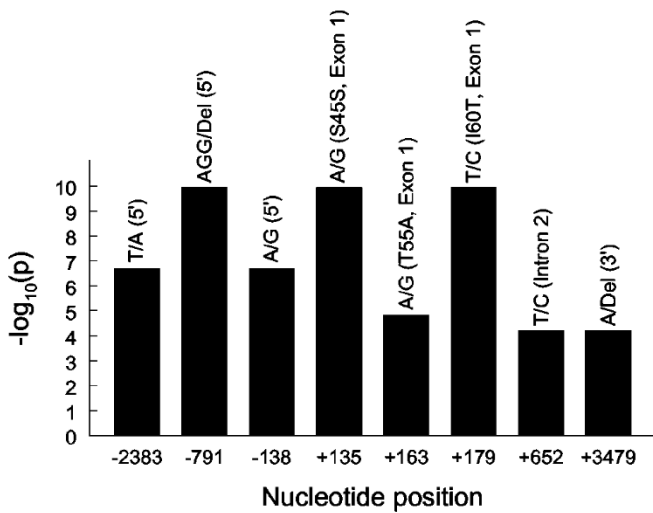
**Figure 1.** Relationship between polymorphisms in the *Tas1r3* region and saccharin preferences in six mouse strains (129P3/J, AKR/J, C57BL/6, CAST/Ei, DBA/2J, and SWR/J). The x-axis shows the polymorphism positions relative to the first nucleotide in the putative start codon of *Tas1r3* (indicated by a vertical line). The y-axis shows the negative decimal logarithm of *p* values obtained in one-way ANOVA tests assessing the effect of genotype for each polymorphism on 1.6 mM saccharin preference. A dotted horizontal line shows the threshold of statistical significance ( $p = 0.01$ ). For the eight polymorphisms significantly associated with saccharin preference, nucleotides (high preferring/low preferring) and their positions are shown; for coding regions, corresponding amino acids and their positions are shown in parentheses. At the bottom, asterisks indicate positions of these eight polymorphisms relative to the *Tas1r3* coding and noncoding regions.

**Table 4. Genotypes of 30 mouse strains at eight polymorphic sites in the *Tas1r3* region associated with 1.6 mM saccharin preference (%)**

Inbred strain	Sac pref	-2383	-791	-138	+135	+163	+179	+652	+3479
CAST/Ei	97	T	<b>AGG</b>	A	A	A	T	T	A
FVB/NJ	96	T	<b>AGG</b>	A	A	A	T	T	A
CE/J	96	A	<b>AGG</b>	G	A	A	T	C	Del
SJL/J	93	T	<b>AGG</b>	A	A	A	T	T	A
SWR/J	93	T	<b>AGG</b>	A	A	A	T	T	A
C57BL/6	91	T	<b>AGG</b>	A	A	A	T	T	A
ST/bJ <sup>g</sup>	91	T	<b>AGG</b>	A	A	A	T	T	A
I/LnJ	91	T	<b>AGG</b>	A	A	G	T	C	Del
SM/J	91	T	<b>AGG</b>	A	A	G	T	C	Del
P/J	89	T	<b>AGG</b>	A	A	A	T	T	A
RBF/DnJ	89	T	<b>AGG</b>	A	A	A	T	T	A
C57L/J	88	T	<b>AGG</b>	A	A	A	T	T	A
KK/HJ	83	T	<b>AGG</b>	A	A	A	T	T	A
SEA/GnJ	83	T	<b>AGG</b>	A	A	A	T	T	A
IS/Cam <sup>g</sup>	83	T	<b>AGG</b>	A	A	A	T	T	A
NOD/LtJ	81	T	<b>AGG</b>	A	A	A	T	T	A
SPRET/Ei	77	T	<b>AGG</b>	A	A	A	T	T	A
NZB/B1NJ	74	T	<b>AGG</b>	A	A	A	T	T	A
PL/J	73	A	Del	G	G	G	C	C	Del
RIIIS/J	71	A	Del	G	G	G	C	C	Del
C3H/HeJ	65	A	Del	G	G	G	C	C	Del
CBA/J	65	A	Del	G	G	G	C	C	Del
RF/J	65	A	Del	G	G	A	C	C	Del
BUB/BnJ	60	A	Del	G	G	G	C	C	Del
BALB/cByJ	59	A	Del	G	G	G	C	C	Del
129P3/J	58	A	Del	G	G	G	C	C	Del
A/J	56	A	Del	G	G	G	C	C	Del
DBA/2J	52	A	Del	G	G	G	C	C	Del
LP/J	48	A	Del	G	G	G	C	C	Del
AKR/J	41	A	Del	G	G	G	C	C	Del

Genotypes creating recombinant haplotypes are underlined. Variant sites most significantly associated with saccharin preference are shown in bold.

<sup>g</sup>Saccharin preference data from Lush (1989).



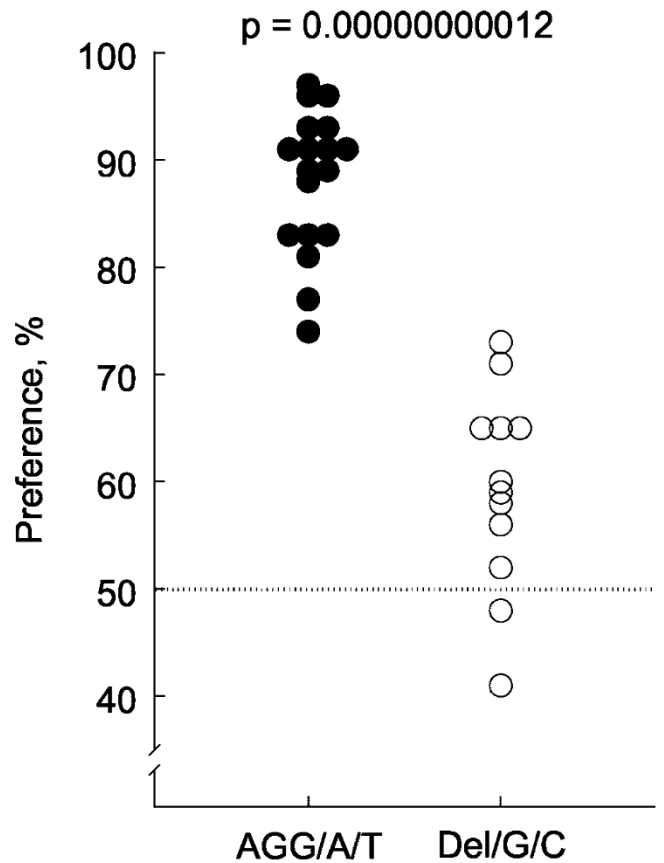
**Figure 2.** Relationship between polymorphisms in the *Tas1r3* region and saccharin preferences in 30 mouse strains. The y-axis shows the negative decimal logarithm of *p* values obtained in *t* tests assessing differences in 1.6 mM saccharin preference between groups of mice with different genotypes at each polymorphic site. Positions, relative to the first nucleotide in the putative start codon of *Tas1r3*, are shown at the bottom. Nucleotide alleles, their locations, and corresponding amino acid substitutions are shown above the bars.

***Tas1r3* gene expression, transcript size, and protein abundance in B6 and 129 mice**

Northern blot analysis identified a single band in both the B6 and 129 strains of mice. There were no differences between strains in the abundance of the transcript, nor was there evidence for alternatively spliced forms of the transcript in either strain (Fig. 4). Likewise, the results of real-time PCR indicated that although the *Tas1r3* gene was expressed in fungiform, circumvallate, and foliate papillae, there were no strain differences in *Tas1r3* gene expression computed relative to either a housekeeping gene (main effect of strain:  $F_{(1,23)} = 0.9$ ;  $p = 0.35$ ) or a taste receptor cell-specific gene (main effect of strain:  $F_{(1,20)} = 2.2$ ;  $p = 0.15$ ) (Fig. 5). As expected from previous research, there was a main effect of tissue-type on *Tas1r3* gene expression. Relative to the housekeeping gene *Tbp*, *Tas1r3* gene expression was lower in fungiform papillae than in circumvallate and foliate papillae ( $F_{(3,23)} = 3.7$ ;  $p = 0.03$ ). Relative to the taste receptor cell-specific gene *Gng13*, *Tas1r3* gene expression was higher in fungiform papillae than in circumvallate and foliate papillae ( $F_{(1,20)} = 5.1$ ;  $p = 0.02$ ). There were no between-strain differences in the expression of the taste cell-specific gene *Gng13* ( $p > 0.05$ ) and no differences between strains in the number of fungiform papillae (A. A. Bachmanov, unpublished observations). Therefore, confounding variables, such as between-strain differences in papillae density or taste receptor cell number, are unlikely to have influenced the measurement of the *Tas1r3* gene expression. The results of immunocytochemistry with an anti T1R3 antibody demonstrated that (1) a subset of taste receptor cells from all types of taste papillae contained T1R3 immunoreactivity, (2) there was little or no T1R3 immunoreactivity in non-taste lingual epithelium, and (3) there were no observable differences in T1R3 immunoreactivity between the B6 and 129 inbred mouse strains (Fig. 6).

**Discussion**

The identification of DNA variant sites within a 6.7 kb region of the *Tas1r3* gene and analysis of genotype–phenotype relationships detected eight polymorphisms significantly associated with saccharin preference among inbred mouse strains. All these vari-

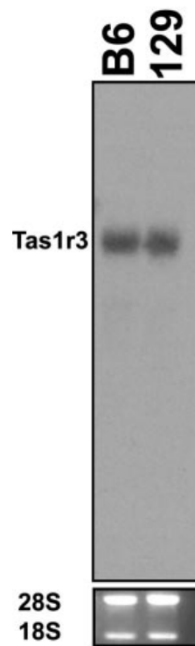


**Figure 3.** Preferences for 1.6 mM saccharin by mice from inbred strains with different *Tas1r3* haplotypes. Closed circles denote means for strains with the nucleotide sequence AGG at nt –791, A at nt +135, and T at nt +179 (AGG/A/T haplotype found in the C57BL/6, C57L/J, CAST/Ei, CE/J, FVB/NJ, I/LnJ, IS/Cam, KK/HUJ, NOD/LtJ, NZB/BINJ, P/J, RBF/DnJ, SEA/GnJ, SJL/J, SM/J, SPRET/Ei, ST/bJ, and SWR/J strains). Open circles show the means for strains with a 3 bp deletion at nt –791, G at nt +135, and C at nt +179 (Del/G/C haplotype found in the 129P3/J, A/J, AKR/J, BALB/cByJ, BUB/BnJ, C3H/HeJ, CBA/J, DBA/2J, LP/J, PL/J, RF/J, and RIIS/J strains). The strains with the AGG/A/T haplotype strongly preferred saccharin (preference score,  $88 \pm 2\%$ ; mean  $\pm$  SE;  $n = 18$ ), whereas strains with the Del/G/C haplotype were indifferent to or only weakly preferred saccharin (preference score,  $59 \pm 3\%$ ;  $n = 12$ ;  $p = 0.0000000012$ ; two-tailed *t* test).

ants are candidates for functional polymorphisms affecting *Tas1r3* biology. The different strength of the genotype–phenotype associations among these polymorphisms provided a hierarchy from most to least feasible candidate functional variant sites.

This association was strongest for three polymorphisms that are, therefore, the most likely candidates for the *Sac<sup>b</sup>* and *Sac<sup>d</sup>* alleles. One of these three polymorphisms, a 3 bp deletion/insertion, is located upstream of the putative initiation codon and could influence *Tas1r3* gene expression. However, we found mixed evidence for this. Although the deletion at this site disrupted a predicted transcription-binding site, its proximity to a repetitive element reduces the likelihood that this deletion has functional significance. Nevertheless, repetitive sequences in upstream gene regions have been shown to affect gene expression (Hung and Penning, 1999). The other key variant is a silent mutation (Ser45Ser) that might affect gene or protein function, for instance, by interfering with correct exon splicing (D’Souza et al., 1999). Using real-time PCR, Northern blot analysis, and immunocytochemistry, we found that *Tas1r3* gene expression, mRNA transcript size, or T1R3 immunoreactivity in taste tissue did not



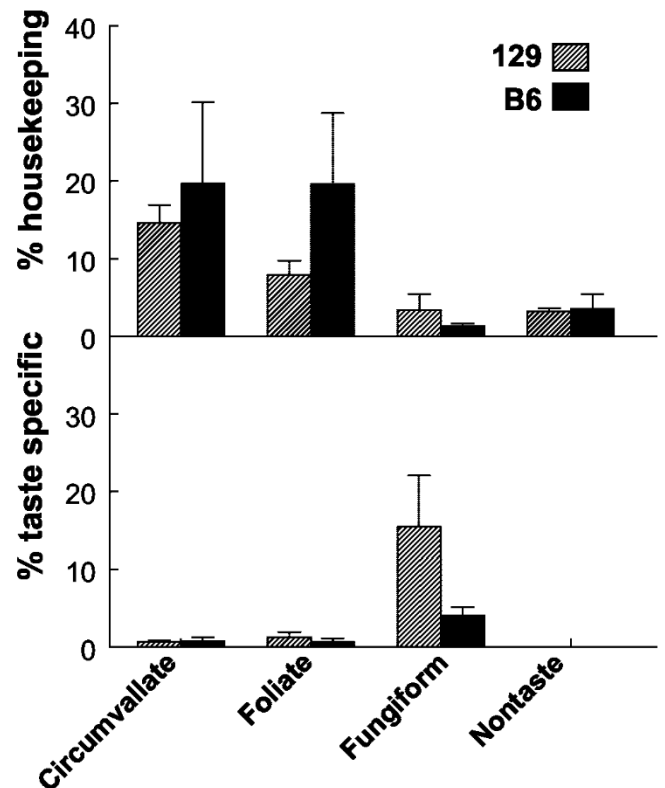


**Figure 4.** Northern blot analysis of *Tas1r3* gene expression in gustatory tissues containing circumvallate and foliate papillae from B6 and 129 mice. Each lane contained 25  $\mu$ g of total RNA. The blot was hybridized with the *Tas1r3* cDNA probe and exposed to film for 4 d. A single major band (7.2 kb) was detected in both mouse strains with similar size and signal intensity. The size of the transcript agrees with previously published Northern blot (Max et al., 2001). Ribosomal RNA (28S and 18S; bottom) is shown as a control for the quality and amount of RNA loaded in each lane.

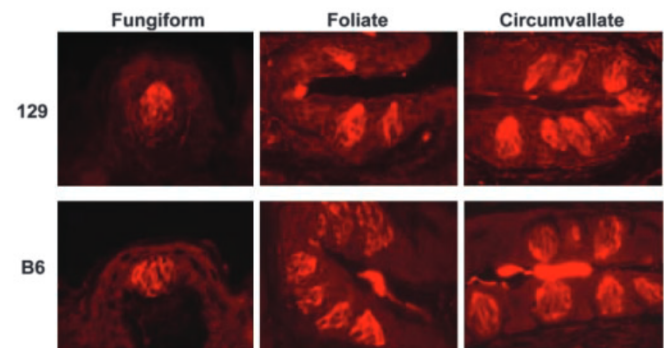
differ between inbred strains with different *Tas1r3* haplotypes. Taken with the results of another study using *in situ* hybridization to measure gene expression (Montmayeur et al., 2001), we conclude that these key polymorphisms are unlikely to act by blocking gene expression in taste tissue. We cannot exclude other effects of these polymorphisms on gene expression, however, such as those on developmental or tissue-specific regulation.

The third key variant resulted in the isoleucine-to-threonine substitution at position 60, in the extracellular N terminus of the predicted T1R3 protein. Modeling of the T1R3 protein using the structure of the related mGluR1 receptor as a prototype has suggested that the Ile60Thr change interferes with an N-linked glycosylation site, which could affect dimerization of the receptor (Max et al., 2001); however, this was not confirmed in a coimmunoprecipitation experiment (Nelson et al., 2002). This type of polymorphism could also affect ligand binding (Rana et al., 2001).

Because these three most likely candidate functional polymorphisms have the same genotype–phenotype associations, we cannot prioritize them. It is possible that the *Sac* alleles are complex, and all three variant sites are needed to affect the phenotype. Such complex alleles have been described in other genes (Phillips, 1999; Drysdale et al., 2000). Moreover, it is possible that there are more than two alleles of the *Sac* gene among the mouse strains examined, so that different polymorphic sites can contribute to different alleles. In fact, we cannot rule out a contribution of any of the eight variants that were significantly associated with saccharin preference, and each one should be considered a candidate functional polymorphism. Consistent with this, one sequence variant resulted in a threonine-to-alanine substitution at position 55, in the predicted extracellular N terminus of the T1R3 protein. Likewise, the nt –2383 SNP affected a predicted transcription



**Figure 5.** *Tas1r3* gene expression as measured by real-time PCR, presented relative to expression of a housekeeping gene (*Tbp*; a TATA box-binding protein) or a taste receptor cell-specific gene (*Gng13*; a G-protein involved in taste receptor signaling). Normalized gene expression was computed using the following formula (percentage of control gene =  $2^{(CT_{control} - CT_{Tas1r3})} \times 100$ ), where *CT* is the PCR cycle number in which the fluorescent product was first detected. Ten samples were eliminated because there was little or no control gene expression (5 samples from nontaste lingual epithelium, 2 samples from circumvallate papillae, 2 samples from fungiform papillae, and 1 sample from foliate papillae). Because the non-taste epithelium does not contain detectable levels of the taste receptor cell-specific gene *Gng13*, there are no values for expression of the *Tas1r3* gene for this category. B6 and 129 mice have similar amounts of *Tas1r3* mRNA in taste papillae relative to either a housekeeping gene or a gene specific to taste receptor cells. Values are presented as means  $\pm$  SE.



**Figure 6.** T1R3 immunoreactivity in taste tissue of B6 and 129 mice. Longitudinal sections from fungiform, foliate, and circumvallate papillae were immunostained with rabbit antiserum directed against a synthetic peptide from the N-terminal domain of T1R3, with a Cy3-conjugated anti-rabbit secondary antibody. No differences between the two mouse strains were apparent in expression levels of the receptor in any sections examined. Magnification, 250 $\times$ .

factor-binding site, and the nt +652 SNP was located in an intron, in which functional polymorphisms can also exist (Fridman et al., 2000). Overall, although nonsynonymous variants are ob-

vious candidates for functional polymorphisms, those in non-coding regions or silent mutations may also have significant biological function.

Positional cloning of quantitative trait loci (QTL), and especially identification of functional polymorphisms defining QTL alleles, is an evolving field. This study demonstrates an approach to examine key sequence variants within a positionally cloned QTL based on genotype–phenotype associations in a large number of genealogically diverse inbred mouse strains. Although special care was taken to select genealogically diverse strains, the results of the Mouse Genome Sequencing Consortium suggest that inbred strains of mice are a mosaic of only a few ancestral mouse haplotypes (Wade et al., 2002) and that blocks of chromosomal segments identical by descent are, on average, ~1 Mb. The results of the current analysis partially confirm this observation. However, several of the inbred strains had mosaic haplotypes that were much smaller than 1 Mb, sometimes spanning only a few hundred base pairs. Therefore, methods of gene and allele identification that assume long stretches of identity by descent among inbred strains (Hitzemann et al., 2002) should be applied with caution.

Our results suggest that allelic variation at the *Tas1r3* gene explains 78% of the strain variation in saccharin preference. However, there was a substantial amount of genetic variation among the inbred mouse strains not attributed to these polymorphisms, which is consistent with evidence for the multigenic determination of taste responses to sweeteners in mice (Pelz et al., 1973; Stockton and Whitney, 1974; Ramirez and Fuller, 1976; Bachmanov et al., 1996a, 1997; Inoue et al., 2004).

In conclusion, we have identified eight polymorphisms in the *Tas1r3* region significantly associated with saccharin preference; three of these are the most likely candidates for functional polymorphisms. These eight variant sites include two protein-coding variants described previously (Bachmanov et al., 2001; Kitagawa et al., 2001; Max et al., 2001; Montmayeur et al., 2001; Nelson et al., 2001; Sainz et al., 2001) and six variants that do not affect predicted T1R3 protein sequence. Here, we presented data for future studies directed to experimentally confirm the function of these polymorphisms and highlighted some of the difficulties of identifying specific DNA sequence variants that underlie QTL.

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