

No Relationship between Sequence Variation in Protein Coding Regions of the *Tas1r3* Gene and Saccharin Preference in Rats

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Abstract

Nearly all mammalian species like sweet-tasting foods and drinks, but there are differences in the degree of 'sweet tooth' both between species and among individuals of the same species. Some individual differences can be explained by genetic variability. Polymorphisms in a sweet taste receptor (*Tas1r3*) account for a large fraction of the differences in consumption of sweet solutions among inbred mouse strains. We wondered whether mice and rats share the same *Tas1r3* alleles, and whether this gene might explain the large difference in saccharin preference among rats. We conducted three experiments to test this. We examined DNA sequence differences in the *Tas1r3* gene among rats that differed in their consumption of saccharin in two-bottle choice tests. The animals tested were from an outbred strain (Sprague–Dawley; experiment 1), selectively bred to be high- or low-saccharin consumers (HiS and LoS; experiment 2), or from inbred strains with established differences in saccharin preference (FH/Wjd and ACI; experiment 3). Although there was considerable variation in saccharin preference among the rats there was no variation in the protein-coding regions of the *Tas1r3* gene. DNA variants in intronic regions were detected in 1 (of 12) outbred rat with lower-than-average saccharin preference and in the ACI inbred strain, which also has a lower saccharin preference than the FH/Wjd inbred partner strain. Possible effects of these intronic nucleotide variants on *Tas1r3* gene expression or the presence of T1R3 protein in taste papillae were evaluated in the ACI and FH/Wjd strains. Based upon the results of these studies, we conclude that polymorphisms in the protein-coding regions of the sweet receptor gene *Tas1r3* are uncommon and do not account for individual differences in saccharin preference for these strains of rats. DNA variants in intron 4 and 5 are more common but appear to be innocuous.

Introduction

Most animals prefer sugars that humans perceive as sweet, and avoid substances that humans perceive as bitter. The acceptance of sweet tastes and the rejection of bitter tastes is present at birth in humans and other primates (Steiner, 1979; Steiner *et al.*, 2001). This observation suggests that taste preferences are innate. Differences exist in the degree to which individual humans (and members of other species) detect sweet or bitter substances (Blakeslee and Salmon, 1935), and an early hypothesis suggested that variability in sweet preferences was due to inherited differences in the 'sensory apparatus' (Nachman, 1959). This hypothesis has recently received support. Three members of a family of genes were discovered (*Tas1r1-3*), and *in vivo* and *in vitro* experiments have suggested that two members of this family

(*Tas1r2* and *Tas1r3*) dimerize and form a sweet receptor (Nelson *et al.*, 2001; Li *et al.*, 2002). Alleles of the *Tas1r3* gene account for much of the difference among inbred mouse strains in saccharin intake and preference (Bachmanov *et al.*, 2001a,b; Kitagawa *et al.*, 2001; Max *et al.*, 2001; Montmayeur *et al.*, 2001; Nelson *et al.*, 2001; Sainz *et al.*, 2001; Inoue *et al.*, 2004; Reed *et al.*, 2004).

The observation that genetic variation in a sweet receptor gene influences sweet preference in mice has a parallel in humans for bitter perception. In most human populations tested, 10–30% of people do not perceive the bitterness of a class of chemical compounds (sometimes called nontasters), whereas other people find them extremely bitter (tasters; Mourant *et al.*, 1976; Guo and Reed, 2001). Because both

tasters and nontasters are also found within some closely related non-human primate species, investigators speculated that the alleles that produce nontasters might be shared between humans and some primates (Fisher, 1939). This idea, that specific alleles of taste genes might be preserved across species, led us to wonder whether the allelic variation found in the mouse sweet taste receptor gene *Tas1r3* might also be present in a related species (rats) and explain the large variation among individuals and rat strains in saccharin preference and intake (Nachman, 1959; Vartiainen, 1967; Overstreet *et al.*, 1993; Dess and Minor, 1996; Giza *et al.*, 1996). To test this hypothesis, we conducted three experiments. In all experiments, we examined DNA sequence differences in the *Tas1r3* gene among rats that differed in their intake and preference for saccharin. The studies used rats with diverse genetic histories: in experiment 1, the subjects were genetically heterogeneous, outbred Sprague–Dawley rats, in experiment 2, they were partially inbred strains that have been selectively bred for high or low saccharin intake (Dess and Minor, 1996), and in experiment 3, they were inbred strains that were known to differ in saccharin intake (Overstreet *et al.*, 1999). When DNA sequence differences were found, they were evaluated in several ways to determine whether the polymorphism might influence gene function and saccharin preference. First, the DNA variants in the rat were aligned against the mouse sequence to determine whether mice with high or low saccharin preferences were allelic in the same locations as the rat. Second, *in silico* methods were used to determine whether the differences in DNA sequence resulted in a predicted change in amino acid, a prematurely truncated protein or would interfere with intron–exon splicing. Finally, because DNA sequence variation outside protein coding regions might influence the amount or distribution of *Tas1r3* mRNA, tissues from four regions of the rat tongue (fungiform, foliate, circumvallate papillae and non-taste epithelium) were compared among rats with *Tas1r3* DNA sequence variation to assess their influence on gene expression and the presence of T1R3 protein.

Materials and methods

Obtaining the rat genomic sequence for *Tas1r3*

When this study began, neither the rat cDNA nor the genomic sequence for *Tas1r3* was known and therefore had to be determined. One genomic DNA sample from the Sprague–Dawley strain (Clontech, Palo Alto, CA) was purchased and served as the reference sequence. Primers were designed based upon the mouse *Tas1r3* sequence (Table 1) and amplified using the following conditions: 94°C, 2 min denaturation; 94°C 45 s, 56–66°C 45 s, 72°C 2 min for 35 cycles; 72°C 7 min extension. All DNA sequencing was done by the University of Pennsylvania Sequencing Core Facility. The obtained sequences were aligned with the mouse and human genomic DNA, as well as the rat cDNA, and the pre-

Table 1 Primers used to amplify genomic fragments of rat *Tas1r3*

Name	Forward	Reverse	Annealing temperature (°C)
RSac1	GGGTAGAAGCCAGTG CTAGCTAAGCG	CTGTCCCAATACCTATA CATGCCAG	60
RSac4	CCCAGTGACCGGG TGCAGCT	GGCACCTGGACATG GAGCCT	66
RSac5	AGGATCCTGAGGAGC TTGAG	TGCAACAGTGTCA CAACGGC	60
RSac7	CCTGCACACACCTTC TATTG		56
RSac12		TAGGGCTGGGTTC CTAAAG	56

dicted amino acid sequence was compared with the mouse (Accession #AF389853) and human (Accession #AC026283; Figure 1). During the project, the rat cDNA sequence became available (Accession #AY032620). This allowed us to check against the sequence we obtained (a match) and to compare the intron and exon structure with the new genomic DNA sequence we obtained.

Animals and saccharin intake

In experiment 1, 12 male CD™IGS (Sprague–Dawley) rats ~40 days old were purchased from Charles River (Wilmington, MA) and acclimated to laboratory conditions for 4 weeks. The daily water and saccharin intakes were measured using a two-bottle choice test, with water as one choice and 3.7 mM sodium saccharin (Sigma Chemical Corp., St Louis, MO) as the second choice. This test was conducted for four consecutive days. The position of the bottles was reversed daily to control for the preference of some rats for solutions on a particular side. Total intakes were averaged for each solution and expressed as the number of milliliters consumed per day. A preference score was computed by dividing the amount of saccharin consumed per day by the total fluid consumed per day. After this measurement was complete, small pieces of tail were obtained from each rat for DNA extraction. Experiment 1 was conducted in the laboratory of Michael G. Tordoff, at the Monell Chemical Senses Center.

In experiment 2, a second group of rats was tested from strains selectively bred to be either high-saccharin (HiS) or low-saccharin (LoS) drinkers (Dess and Minor, 1996). Both the selective breeding and the behavioral testing were conducted in the laboratory of Nancy K. Dess at Occidental College. The selective breeding project was initiated upon the incidental discovery of a male rat that was averse to saccharin solution. This male and an avid saccharin-drinking male were each mated with different pairs of females with average saccharin preference to create the Occidental Low-Saccharin (LoS) and High-Saccharin (HiS) lines. Rats in each litter with the most extreme phenotypes were selected

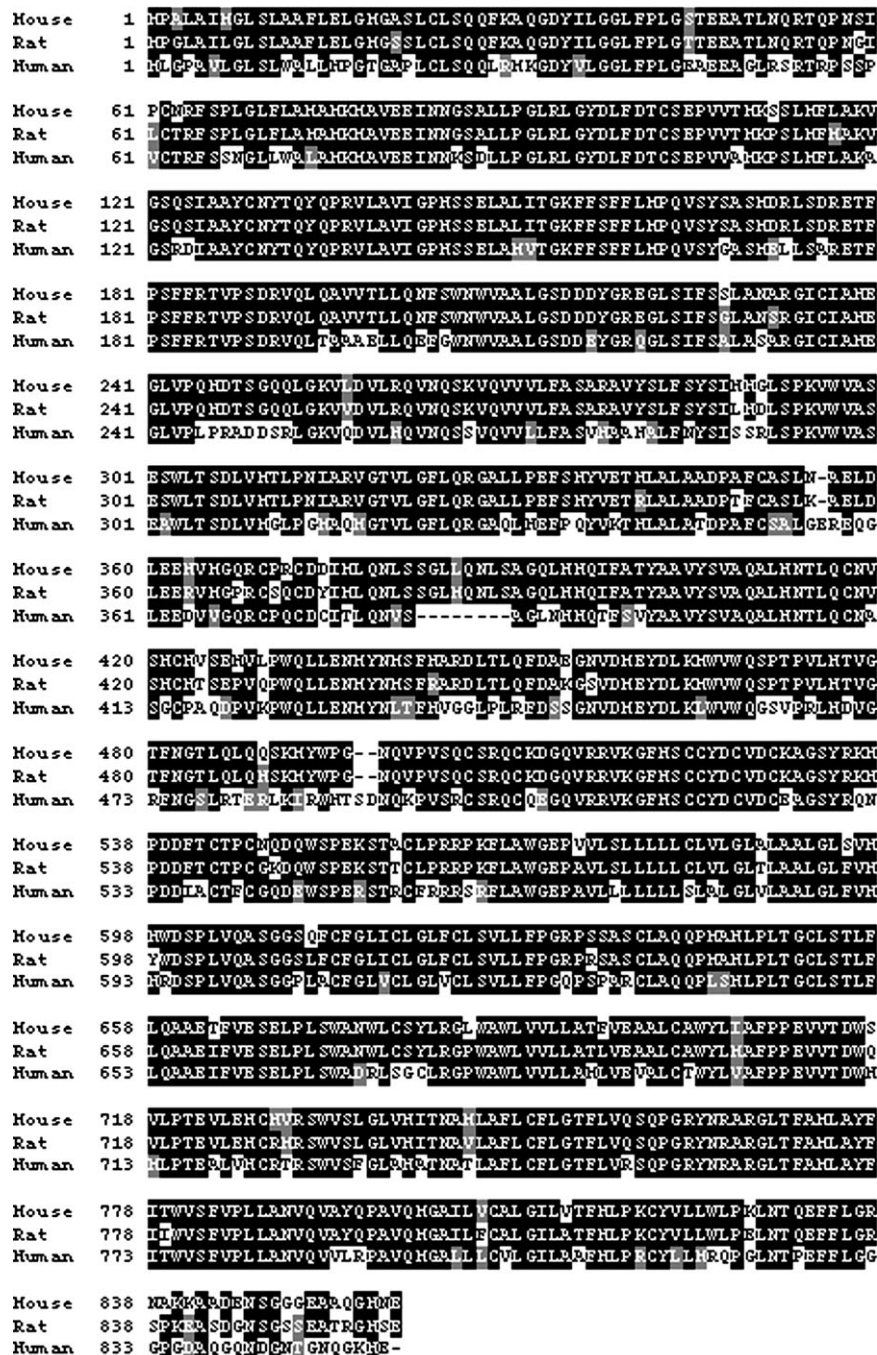


Figure 1 Alignment of predicted mouse (strain C57BL/6ByJ), rat (strains Sprague–Dawley, HiS, LoS, ACI and FH/Wjd strains), and human VTR3 protein. The amino acid sequence was aligned with ClustalW and plotted with BOXSHADE. Black boxes indicate regions of identity, and gray boxes indicate conservative amino acid substitutions. In mice, higher or lower saccharin preferences are associated with a polymorphism at aa position 60 (isoleucine to threonine).

for breeding, excluding rats with extreme baseline water intake. No matings occurred between rats that were quarter-sibs or closer in genetic relatedness, and ~15% of the rats bred every four or five generations are founders from the original stock (formerly Holtzman, now Harlan Sprague–Dawley Holtzman). The rats used in the present study were

from five litters (two HiS, three LoS) in generations 19 and 20. They were phenotyped with a 2 day water-only baseline period followed by a 24 h two-bottle choice between water and 4.5 mM saccharin solution. Preference scores were computed by dividing the amount of saccharin consumed by the total amount of fluid consumed. After the behavioral testing

was completed, the tail of every subject was clipped for DNA extraction.

In experiment 3, FH/Wjd and ACI rats were bred at the Monell Chemical Senses Center from groups of founder rats provided by Drs Amir Rezvani and David Overstreet at the Center for Alcohol Studies, University of North Carolina–Chapel Hill. (The FH/Wjd inbred strain is sometimes referred to as the Fawn Hooded strain.) Eight male and female rats from the inbred FH/Wjd strain and eight male and eight female rats from the ACI strain were offered ascending concentrations of saccharin to drink, with each solution offered for 2 days. Group means were compared by ANOVA, using sex, strain and concentration as factors. The 3.2 mM saccharin solution corresponds most closely to the concentrations used in experiments 1 and 2. Genomic DNA from an FH/Wjd and an ACI rat were obtained for DNA sequencing. Because the FH/Wjd and ACI strains are inbred, all rats should be genetically identical, and therefore only one DNA sample per strain was needed to detect genetic differences within the *Tas1r3* gene. Experiment 3 was conducted in the laboratory of Michael G. Tordoff at the Monell Chemical Senses Center.

Note that due to procedural differences across laboratories, slightly different concentrations were used to test the rats' preference for saccharin in experiments 1–3 (3.7, 4.5 and 3.2 mM respectively). These concentrations were chosen because they were clearly detectable; they are classically considered to fall close to but slightly below the saccharin concentration most preferred by rats (Young and Schulte, 1963). Although there is some evidence that sweetener intake of mice is under differential genetic control at threshold versus suprathreshold concentrations (Bachmanov *et al.*, 1997), this is unlikely to be a factor for the saccharin concentrations used here.

Sequencing of rat genomic DNA samples

Genomic DNA was extracted from tails (Gentra Systems, Minneapolis, MN), and primers were designed from the mouse sequence and then modified, based on amplified rat sequence, to overlap the gene: primer sets were selected to amplify large fragments of rat *Tas1r3* (RSac5F and RSac5R, 1100 bp; RSac4F and RSac4R, 1142 bp; RSac1F and RSac1R, 1773 bp; RSac7F and RSac12R, 3236 bp), and then nested primers were used to close sequencing gaps (Table 1). To reduce the likelihood that errors in the fidelity of Taq polymerase could account for sequence differences among strains, we used a Taq polymerase from *Pyrococcus furiosus*, which has the lowest error rate of known thermophilic DNA polymerases. Polymerase chain reaction (PCR) products were purified (Qiagen, Valencia, CA) and sequenced. Strands were aligned using Sequencher (Genecodes, Ann Arbor, MI). Nucleotide bases were compared and polymorphisms identified. When polymorphisms were found, they were confirmed by additional PCR reactions followed by sequencing.

In silico analysis of DNA sequence variants

Using the reference intron–exon structure of *Tas1r3*, alterations of DNA sequence variants were assessed for predicted protein-coding changes, stop codons or splice junction variants (Discovery Gene Studio version 1.5, San Diego CA). Next the polymorphic site and its flanking region from each rat strain was aligned with the homologous region from the *Tas1r3* gene from strains of mice chosen for higher and lower saccharin intake (Reed *et al.*, 2004). The rationale for this procedure is twofold. First, the degree of DNA sequence conserved across species may be commensurate with its degree of biological importance. Second, if identical polymorphisms were found in mice with higher and lower saccharin preference and these polymorphisms segregated with saccharin intake, it would suggest that the rat polymorphisms might also have an influence on saccharin intake. Mouse sequences from the *Tas1r3* gene were obtained by PCR amplification of genomic templates from six inbred strains, three with higher and three with lower saccharin preferences. Both forward and reverse strands were sequenced and aligned with Sequencher (GeneCodes, Ann Arbor, MI), and rat and mouse sequences were aligned with ClustalW (Chenna *et al.*, 2003), and plotted with BOX-SHADE (version 3.21) (Figure 4). Percent identity among the aligned sequences was computed using the algorithm *bl2seq* (Tatusova and Madden, 1999). These mouse *Tas1r3* DNA sequences have been published previously (Reed *et al.*, 2004).

Real-time PCR

It is possible that intronic sequence variants might lead to markedly reduced gene expression that in turn might result in fewer receptors and an altered taste preference. To determine whether *Tas1r3* gene expression differed among inbred strains of rats with different genotypes, we compared the relative abundance of *Tas1r3* gene expression in fungiform, foliate, circumvallate and non-taste epithelium [corrected for gustducin gene expression in one analysis and for a ubiquitously expressed control gene expression (*B2m*) in a second analysis]. Lingual tissues used for real-time PCR and immunocytochemistry (below) were dissected as previously described (Huang *et al.*, 1999). The original mouse protocol was modified for rats by raising the volume of proteinase enzyme injected underneath the lingual epithelium (6 μ l), and incubating the tongue for 60 min. Total RNA was extracted from the lingual tissues of four rats (two adult male rats from each inbred strain) using the Stratagene Absolutely RNA™ Microprep kit (La Jolla, CA). We could not study SD rat 11 (which also had intronic DNA sequence variants) because the tissue from this rat was not available. First strand synthesis of cDNA was conducted following the directions of the manufacturer (Invitrogen, SuperScript™ III). Transcribed cDNA (1 μ l/well) was transferred to a PCR 96-well optical reaction plate. Each well was supplemented with

Table 2 Experiment 1, saccharin intake of outbred Sprague-Dawley rats

Rat no.	Body weight (g)	Water intake (ml)	3.7 mM saccharin intake (ml)	Preference (%)	Genotype
SD 6	258	21	16	43	reference
SD 8	229	16	74	83	reference
SD 11	236	4	43	91	three variants ^a
SD 10	231	5	68	94	reference
SD 12	292	3	81	96	reference
SD 2	213	2	57	97	reference
SD 3	236	2	66	97	reference
SD 4	266	2	53	97	reference
SD 7	210	2	58	97	reference
SD 5	243	2	97	98	reference
SD 9	234	1	81	98	reference
SD 1	251	1	122	99	reference

Water intake and saccharin intake are given as the average amount consumed per day during a 4 day test.

^aSee Table 5 for a description of polymorphisms. Reference = identical to previously published sequence (Accession # AY032620).

predeveloped TaqMan assay reagents (primer and fluorescent probe, for location of probe, see Figure 3) and Universal PCR Master Mix™ to a final volume of 50 μ 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 s and 60°C for 60 s. DNA amplification was performed in an ABI PRISM 7000 Sequence Detection System (Applied Biosystems), detecting fluorescence (500–560 nm) after laser light excitation. PCR cycle threshold (CT) values represent the time at which the emitted fluorescence increased above threshold. The *Tas1r3* primers spanned the intron between exons 4 and 5 (Accession # NM_130818, assay ID # Rn_00590759; Figure 3). *Tas1r3* gene expression was normalized as previously described (Livak and Schmittgen, 2001) to a gene that is selectively expressed in taste receptor cells (gustducin, *Gnat3*; Accession # NM_173139) or an ubiquitously expressed control gene (*B2m*; Accession # NM_012512) relative gene expression was calculated using non-taste tongue epithelium was used as a calibrator tissue. Few animals were available for study and statistical analysis of the strain differences were therefore not possible.

Immunocytochemistry

Polyclonal antisera against a hemocyanin-conjugated T1R3 peptide (aa 45–62) were raised in rabbits (see Figure 3 for the location of aa sequence). Frozen sections of rat 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 h at room

Table 3 Experiment 2, rats selectively bred for high (HiS) and low (LoS) saccharin intake

Rat no. and group	Body weight (g)	Water intake (ml)	4.5 mM saccharin intake (ml)	Preference (%)	Genotype
5 LoS	319	6	58	91	reference
6 LoS	288	8	60	88	reference
1 LoS	523	13	63	83	reference
4 LoS	450	1	71	99	reference
Mean \pm SD	395 \pm 111	7 \pm 5	63 \pm 6	90 \pm 7	
8 HiS	277	1	212	100	reference
2 HiS	485	1	230	100	reference
3 HiS	529	1	234	100	reference
7 HiS	283	2	243	99	reference
Mean \pm SD	394 \pm 132	1 \pm 0.4	230 \pm 13	100 \pm 0	

Water intake and saccharin intake are given as the total amount consumed during a 2 day test.

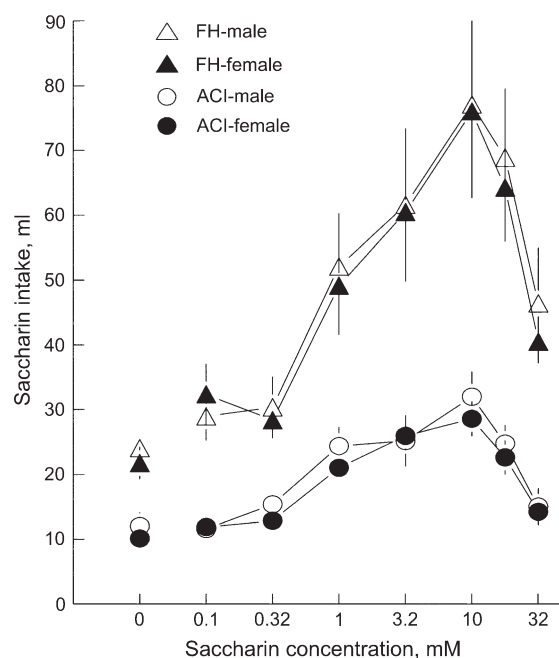


Figure 2 Mean (\pm SEM) saccharin intakes of male and female FH/Wjd and ACI rats. Intakes are daily averages of 48 h tests. Water intake for all groups is shown as 0 mM saccharin. Water intake during tests with saccharin available was minimal (not shown).

temperature and then incubated for 16 h 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 μ M. Preimmune serum did not show any immunoreactivity.

Table 4 Experiment 3, 3.2 mM saccharin intake of two inbred rat strains

Strain	Water intake (ml)	Saccharin intake (ml)	Preference (%)	Genotype
ACI	3.6	25 ± 4	83 ± 9	two variants*
FH/Wjd	4.8	62 ± 12	90 ± 5	reference

Values presented are averages from eight male rats from each inbred strain offered water and 3.2 mM saccharin to drink.

See Table 5 for a description of the polymorphism and Figure 2 for details concerning saccharin consumption.

Table 5 Description of polymorphisms within the rat *Tas1r3* gene

Experiment	Rat	Intron		
		4	5	5
		+1873 nt	+2130 nt	+2164 nt
1	SD #11	G/G to G/A	G/G to G/A	C/C to C/T
3	ACI	G/G to A/A	G/G to A/A	reference (C/C)

Nucleotides are numbered with A in the putative methionine start codon as +1.

Results

Construction of the rat sequence of *Tas1r3*

We obtained 3912 bp of DNA sequence that composed and flanked the rat *Tas1r3* gene. The rat gene is highly homologous at the amino acid level to the mouse and human gene (Figure 1). Similar to the mouse gene, the rat gene has six exons, interrupted by small introns. We found several polymorphisms, as described below, but none within the protein coding regions. Specifically, the I60T polymorphism in the mouse protein coding region of the *Tas1r3* gene associated with saccharin preference in mice was the 'taster' form of the allele (i.e. isoleucine at aa position 60) in all rats studied.

Experiment 1

Sprague-Dawley rats. On average, rats drank 5.1 ± 6.3 ml (mean ± SD) of water and 66.7 ± 26.8 ml of 3.7 mM saccharin per day (Table 2). With one exception (rat 6), all rats drank more saccharin solution than water (i.e. percent preference scores were >50%). Some rats drank prodigious amounts of saccharin, and one rat (rat 1) consumed, on average, almost half of its body weight in saccharin solution daily. Eleven of the 12 Sprague-Dawley rats had a sequence of the *Tas1r3* gene that was the same as the reference sequence (Table 2). Three heterozygous sequence variants were detected within the *Tas1r3* gene for rat 11 (Table 5). This rat drank ~20 ml less than the group average but consumed >90% of its fluid as saccharin.

Experiment 2

Rats selectively bred to differ in saccharin intake. Rats of the LoS strain had low intakes of saccharin and drank between 58 and 63 ml/day, whereas rats of the HiS strain drank between 212 and 243 ml/day. All rats showed a preference for saccharin, but the LoS preference was lower compared with the HiS strain (Mann-Whitney $U = 0.50$, $P = 0.026$; Table 3). However, all rats, bred from either the LoS or the HiS line, had the same *Tas1r3* genotypes as the reference sequence (Table 3).

Experiment 3

ACI. FH/Wjd rats differ in sequence of the *Tas1r3* gene. Saccharin intake was significantly higher in the FH/Wjd than in the ACI rats, $F(1,30) = 41.56$, $P < 0.000001$ (Fig. 2), which confirms earlier observations that the FH/Wjd strain is a saccharin-preferring inbred strain. The FH/Wjd strain had the reference sequence and the ACI strain had two sequence variants (Tables 4 and 5; Figure 3): a substitution of G to A at nt 1873 was found in intron 4, and a G to A substitution at nt 2130 was found in intron 5. Neither of these polymorphisms was within the splice junction and would not therefore be predicted to interfere with the creation of the correct exon order and length (Figure 3).

Comparison of the intronic rat DNA sequence variants with DNA sequences from inbred mouse strains with high and low saccharin preference

The DNA sequence flanking the rat intronic variants was aligned with homologous sequence of mice selected for high and low saccharin intake (Bachmanov *et al.*, 2001a,b; Reed *et al.*, 2004) to determine whether the nucleotides were conserved and whether a pattern of polymorphism was shared between rats and mice with similar saccharin preference (Figure 4). The homology between rat and mouse sequence from intron 4 and 5 was high (>80%), and was higher for intron 4 (75/82 identity; 91%) than for intron 5 (140/172; 81%). The intron 4 rat polymorphism (G to A at position +1873) was a G in all mice regardless of saccharin preference. For the two polymorphisms in intron 5, the first polymorphism (G to A at +2130) was not conserved between mouse and rats. This nucleotide was deleted in all mouse strains studied. The second polymorphism in intron 5 was a C at position +2164 for both the FH/Wjd and ACI rat strains, whereas the SD rat 11 and the six mouse strains shared a T at this position. Of the three *Tas1r3* polymorphisms identified in the rats, none shared a common pattern of polymorphism with mice strains selected for high and low saccharin preference.

Tas1r3 gene expression

Gene expression in the foliate papillae was more abundant relative to the fungiform or circumvallate papillae, and was

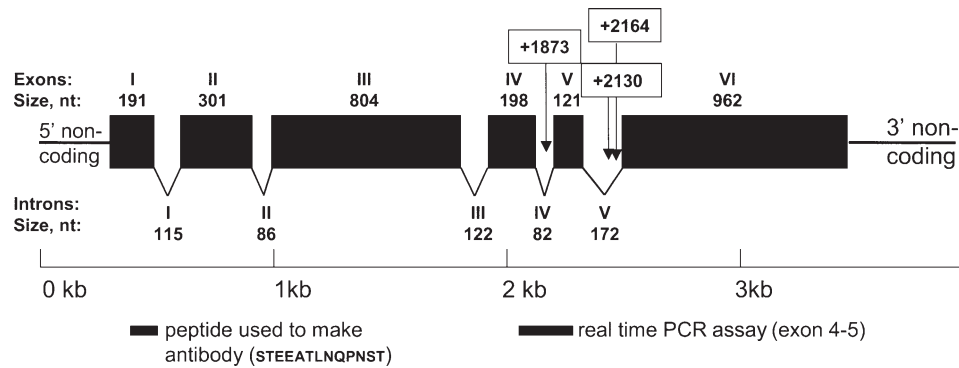


Figure 3 Intron-exon structure of the rat *Tas1r3* gene. Above the gene structure, exons are labeled in Roman numerals, with their size (in nucleotides) shown underneath. Below the gene structure, introns are labeled in Roman numerals, and their size is shown underneath. The nucleotide positions are numbered such that the first nucleotide of the putative methionine initiation codon is +1. Polymorphisms are denoted by the changed nucleotide position and are shown in open boxes. Intron and exon structure was determined by aligning cDNA with genomic DNA. Filled boxes underneath the gene structure show the locations of the real time PCR probe and the peptide used to produce the antibody.

Tas1r3 Intron 4

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FH/Wjd      GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAAC-----CAATGC TAAGCTAGCG G AG-CCCTACCCAGGCTCCATGTCCAG
ACI         GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAAC-----CAATGC TAAGCTAGCG A AG-CCCTACCCAGGCTCCATGTCCAG
SD#11      GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAAC-----CAATGC TAAGCTAGCG A AG-CCCTACCCAGGCTCCATGTCCAG
B6mouse    GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAACCTTGTCCGCTTTGGGGCAATGC TAAGCTAGCG G AGGCTACCCAGGCTCCATGTCCAG
129_DBA_SWR_AKRmouse GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAACCTTGTCCGCTTTGGGGCAATGC TAAGCTAGCG G AGGCTACCCAGGCTCCATGTCCAG
CASTmouse  GTAAGCACTAGACAGCCAAAAGCCATGGTGGCTGAAACCTTGTCCGCTTTGGGGCAATGC TAAGCTAGCG G -GGCTACCCAGGCTCCATGTCCAG

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Tas1r3 Intron 5

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FH/Wjd      CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAATTACTGTGGGG G CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTT---CA C CACTACAG
ACI         CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAATTACTGTGGGG A CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTT---CA C CACTACAG
SD#11      CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAATTACTGTGGGG A CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTT---CA T CACTACAG
B6mouse    CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAAGTTACTGTGGGG - CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTTCTCTTCCA T CACTACAG
129mouse   CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAAGTTACTGTGGGG - CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTTCTCTTCCA T CACTACAG
CASTmouse  CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAAGTTACTGTGGGG - CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTTCTCTTCCA T CACTACAG
DBA_SWR_AKRmouse CTGGCAGGCCCCCTCCGCCAGGGCTAATGTTGAGGCAAGTTACTGTGGGG - CTGGCTGGG GAGTTCGTCTCCCATCACCAGCCCCACATTCGCTTTCTTTCTCTTCCA T CACTACAG

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Figure 4 Alignment of the *Tas1r3* sequence (introns 4 and 5) from the FH/Wjd and ACI inbred strains, as well as from SD rat 11, and six inbred strains of mice with high (C57BL/6J, SWR/J, CAST/Ei) and low (129P3/J, DBA/2J, AKR/J) saccharin preference. All other rats from experiments 1 and 2 had the same DNA sequence as the rats from the FH/Wjd strain. For the mice, inbred mouse strains with identical DNA sequence are shown as a single sequence. The abbreviations for the mouse strains are as follows, C57BL/6J (B6), SWR/J (SWR), CAST/Ei (CAST), 129P3/J (129), DBA/2J (DBA) and AKR/J (AKR). The rat polymorphic sites (+1873 nt, intron 4; +2130 and +2164, intron 5), are shown in the rectangular boxes.

somewhat higher in the FH/Wjd strain compared with the ACI strain (Figure 5). There was little or no *Tas1r3* gene expression in non-taste epithelium. The pattern of gene expression in tongue tissue is consistent with previous reports of the expression pattern of *Tas1r2* (Hoon *et al.*, 1999), a gene that co-localizes with *Tas1r3* in taste receptor cells (Li *et al.*, 2002). Similar patterns of *Tas1r3* expression were found whether the abundance was expressed relative to gustducin, a taster-receptor cell-specific gene (Huang *et al.*, 1999), or *B2m*, a ubiquitously expressed control gene.

T1R3 immunocytochemistry

The results of immunocytochemistry with an anti-T1R3 antibody demonstrated that (i) a subset of taste receptor cells from all types of taste papillae contained T1R3 immunoreactivity and (ii) there was little or no T1R3 immunoreactivity in non-taste lingual epithelium (Figure 6).

Discussion

Polymorphisms within the *Tas1r3* protein-coding regions make no contribution to saccharin preference in the rats tested. We tested the hypothesis that polymorphisms within the protein-coding regions of the sweet taste receptor gene *Tas1r3* account for individual differences in saccharin intake among rats and, further, that rats with low saccharin preference might have the same alleles that predict amino acid changes in *Tas1r3* as mice with low saccharin preference (Reed *et al.*, 2004). We found no protein-coding sequence variants among the rats tested, and no DNA sequence variation that matched alleles in mice with lower saccharin preference. While sequence variants were found in the introns of the rat *Tas1r3* gene, there was little correspondence between these polymorphisms and behavior. The FH/Wjd rats may have somewhat higher *Tas1r3* gene expression levels compared to the ACI strain. Both the FH/Wjd and ACI strains have

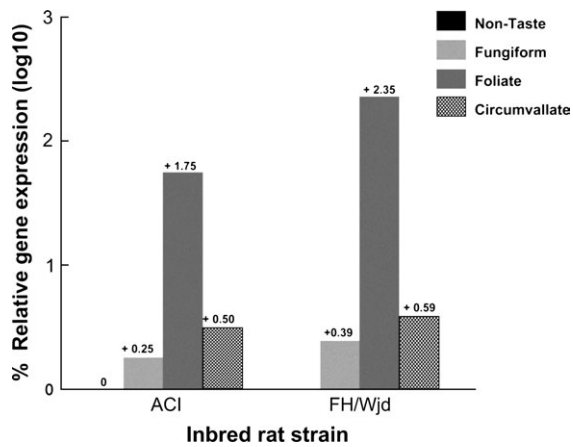


Figure 5 Expression of the *Tas1r3* gene in the fungiform, foliate, circumvallate papillae calibrated to expression in non-taste epithelium of the rat tongue from inbred strains with higher (FH/Wjd) or lower (ACI) saccharin intakes. Relative *Tas1r3* gene expression, either expressed as a proportion of control gene expression (*B2m*, as shown) or gustducin (data not shown) was similar between strains, although the expression in foliate papillae was somewhat higher in the FH/Wjd than the ACI strain.

T1R3 protein in the appropriate lingual tissues as measured by immunocytochemistry. Thus, our results do not support the hypothesis that *Tas1r3* polymorphisms underlie the range of saccharin preferences observed in rats, although there is always the possibility that rare *Tas1r3* polymorphisms may exist in the SD strain (or other outbred strains) or that inbred rat strains not tested here might have alleles in the protein-coding regions of the *Tas1r3* gene.

The *Tas1r3* gene codes for a receptor that is thought to work by binding sugars and other sweeteners in taste receptor cells, and it regulates the intensity of the sensory signal sent to the brain (Bachmanov *et al.*, 1997). This observation suggests that the effect of different alleles of this gene on behavior is limited to the perception of compounds that bind to the receptor such as sucrose and saccharin (Nelson *et al.*, 2001). However, rats selected and bred for high or low saccharin intake have many behavioral differences (e.g. brain self-stimulation, ethanol consumption) in addition to their sweetener preference (Ganchrow *et al.*, 1981; Gosnell and Krahn, 1992; Overstreet *et al.*, 1993; Dess and Minor, 1996; Dess *et al.*, 1998; Kampov-Polevoy *et al.*, 1999; Overstreet *et al.*, 1999; Dess, 2000). Therefore, genes involved in central sensory and hedonic processing, such as those underlying brain reward pathways, rather than in peripheral sensitivity, may account for differences in saccharin preference among rats.

Alternatively, or additionally, allelic differences in other genes contributing to the taste pathway may be involved. In addition to *Tas1r3*, two members of the same receptor gene family (*Tas1r1* and *Tas1r2*) also transduce taste information. *Tas1r1* and *Tas1r2* were discovered several years before *Tas1r3*, and originally *Tas1r1* was thought to be the *Sac* gene (Hoon *et al.*, 1999). Genetic mapping studies discon-

firmed this hypothesis (Li *et al.*, 2001), and later studies have suggested that *T1R1* combines with *T1R3* to transduce information about amino acid rather than sweet taste (Nelson *et al.*, 2001; Li *et al.*, 2002). Therefore, alleles in the *Tas1r1* gene are unlikely to account for differences in saccharin intake. The *Tas1r2* gene is a better candidate gene because its protein product is thought to combine with T1R3 to become a sweet taste receptor. However, although alleles in the *Tas1r2* gene might be involved in individual differences in sweet taste perception in some strains of rats, the search for rat quantitative trait loci for saccharin preference using genome scan methods have not implicated the region containing the *Tas1r2* gene (Murphy *et al.*, 2002). Alleles in bitter receptors or in cell signaling might also influence saccharin intake because these might enhance or diminish the bitter taste of saccharin. Finally, individual differences in central integration of taste information may influence saccharin intake, for instance, rats that prefer water instead of saccharin may perceive saccharin as bitter and salty (Giza *et al.*, 1996).

Polymorphisms in the flanking regulatory sequence may contribute to the difference in saccharin preference. Intronic DNA variants were associated with reduced saccharin intake of the ACI compared with the FH/Wjd strain, and it is possible that these variants have functional consequences on gene expression levels or alternative splicing. The immunocytochemistry and measures of gene expression presented here suggest that there are no large differences between the ACI and FH/Wjd strains in *Tas1r3* gene expression or T1R3 protein abundance in the taste tissues tested. However, the somewhat higher expression of the *Tas1r3* gene in the foliate papillae of the FH/Wjd strain compared with the ACI strain may be worthy of further study. Because the probes used to measure gene expression span exons 4 and 5, it is unlikely that intronic variants in the rats studied cause a disruption in the correct intron–exon splice for these nearby exons. Because the promoter and enhancer of *Tas1r3* are not characterized, we did not look for polymorphisms in the flanking regulatory sequence. The possibility exists that sequence variants among these regions contribute to the difference in saccharin preference among the rats tested.

In addition to the sequencing of candidate genes, two other approaches are worth further consideration to understand genetic variation and sweetness perception in the rat. One approach would be to survey inbred and outbred rat strains for individual or strain differences in response to other sweeteners such as sucralose (Sclafani and Clare, 2004) or fructose (De Francisco and Dess, 1998) and to assess heritability for these traits. Second, when and if high heritability is found, genome scans (a method whereby all genomic regions are assayed for the presence of genes that contribute to behavioral variation) could be a next step in identifying genes involved in sweet taste perception. Several rat genome scans have been conducted using saccharin preference as a trait but the investigators were primarily interested in ethanol

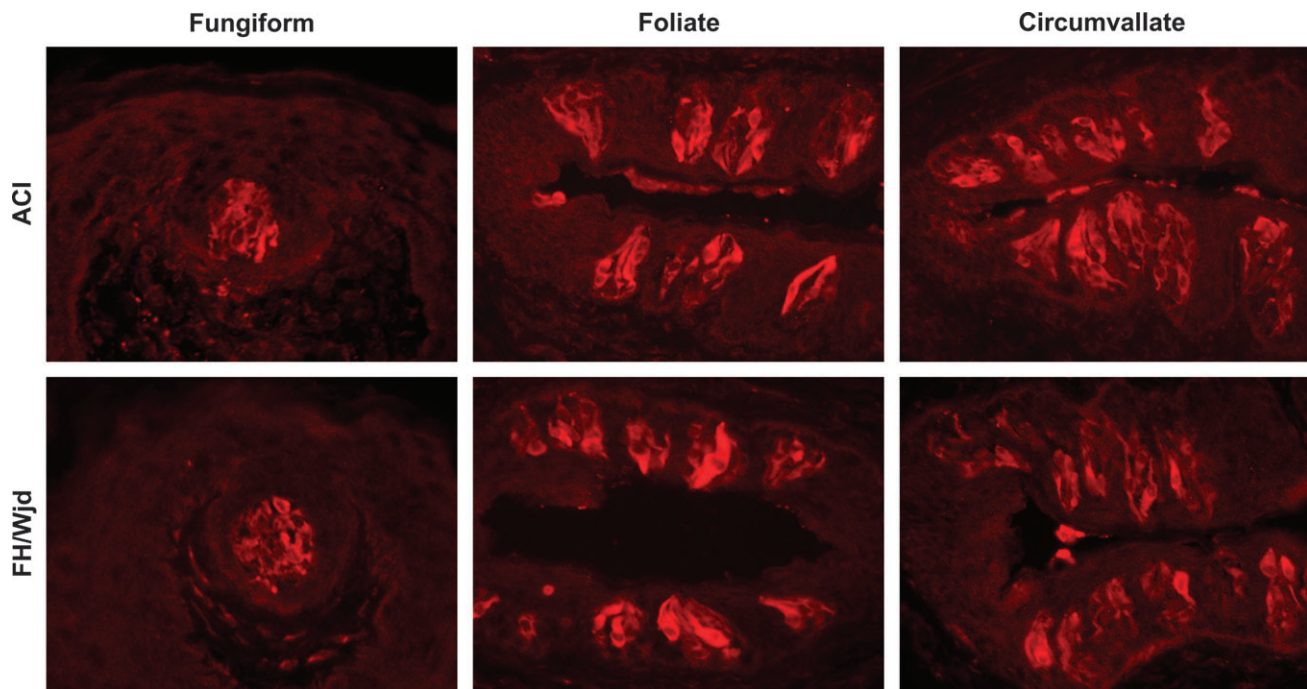


Figure 6 T1R3 immunoreactivity in taste tissue of FH/Wjd and ACI rats. Longitudinal sections from fungiform, foliate, circumvallate and papillae were immunostained with rabbit antiserum directed against a synthetic peptide from the N-terminal domain of T1R3, with a Cy3-conjugated anti-rabbit secondary antibody. No differences in expression levels of the receptor in all sections examined from the two rat strains were apparent. See Figure 3 for location of amino acid sequence used to generate the antibody. ¹Magnification: 250 \times .

and therefore the rats strains selected for study differed in ethanol as well as saccharin preference (Foroud *et al.*, 2002; Terenina-Rigaldie *et al.*, 2003). None of the linkage peaks were near the *Tas1r3* gene. Along with a candidate gene method as reported here, these other approaches are likely to lead to a better understanding of taste and genetics, which in turn may increase our understanding of the mammalian sweet tooth and ultimately have implications for human health.

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