

A *Drosophila* gustatory receptor required for the responses to sucrose, glucose, and maltose identified by mRNA tagging

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In *Drosophila*, detection of tastants is thought to be mediated by members of a family of 68 gustatory receptors (Grs). However, only one receptor, Gr5a, has been associated with a sugar, and it appears to be activated specifically by trehalose. It is unclear whether other sugar receptors are activated by single or multiple sugars. Currently, no Grs are known to colocalize with Gr5a. Such Grs would be candidate sugar receptors because Gr5a-expressing cells function in the responses to attractive tastants. Here we use an "mRNA tagging" approach to identify Gr RNAs that are coexpressed with Gr5a. We found that all seven Grs most related to Gr5a (*Gr64a-f* and *Gr61a*) were expressed in Gr5a-expressing cells, whereas none of the other Grs examined were enriched in these Gr neurons (GRNs). We characterized the role of one Gr5a-related receptor, Gr64a, and found that it was required for the behavioral responses to glucose, sucrose, and maltose. Gr64a was required for GRN function because action potentials induced by these sugars were dependent on expression of Gr64a in GRNs. These data demonstrate that multiple Grs are coexpressed with Gr5a and that *Drosophila* Gr64a is required for the responses to multiple sugars.

gustatory receptor neuron | taste | tip recordings | sugars | behavior

The ability to sense sweetness is fundamental to the survival of animals, ranging from flies to humans, because it facilitates the identification of edible, nutrient-rich sources. In *Drosophila*, detection of sweet, bitter, salt, and pH occurs in gustatory receptor (Gr) neurons (GRNs), rather than neuroepithelial cells, as in mammals (1–3). GRNs are housed in hair-like structures, referred to as sensilla, distributed at the tip of the proboscis (labella), legs, wing margins, and the female genitalia (2).

Drosophila has emerged as an excellent animal model to characterize the sensation of taste. Many studies have focused on the development, distribution, and fine structure of the gustatory organs (2). Moreover, there is recent progress describing the projection patterns of the GRN axons into the brain (4, 5). However, the mechanisms underlying the detection and transduction of tastant-induced stimuli, such as those involved in the sensation of sweetness, are poorly understood.

In mammals the ≈ 45 Grs fall into two families (T1R and T2R). Multiple T2Rs have been shown to function as bitter receptors, leading to the proposal that all T2Rs may be homomeric bitter receptors (6–13). In contrast, the umami response appears to be mediated by a T1R1/T1R3 heteromer (7, 14–16). Surprisingly, two receptors, T1R2 and T1R3, may account for all responses to sugars through distinct combinations of T1R2/T1R3 heteromers and/or T1R2 and T1R3 homomultimers (7, 14–20). In *Drosophila* there is a single family of 68 Grs (21–24), and the proportion devoted to sweet, as opposed to bitter, tastants is unresolved. Currently, only two *Drosophila* Grs have been associated with specific tastants. These include Gr66a, which is required *in vivo* for the avoidance behavior to caffeine (25), and Gr5a, which has been reported to respond specifically to trehalose (26–28).

Expression of *Gr66a* and *Gr5a* reporters indicates that these two Grs are expressed in nonoverlapping subsets of GRNs and appear to define most, but not all, of the GRNs that respond to

avoidance and attractive compounds, respectively (4, 5). At least nine additional Grs are expressed in the labellum, and, based on transgenic reporters, all are coexpressed in subsets of the *Gr66a*-positive GRNs (4, 5). Thus, as in mammals (3), it appears that the majority of Grs participate in the perception of aversive compounds.

In addition to Gr5a, there must be additional sugar-stimulated Grs because Gr5a has been reported to be tuned specifically to trehalose and *Gr5a* mutant flies display normal behavioral and electrophysiology responses to sucrose (26–28). Other sugar receptors would appear to be coexpressed with Gr5a because Gr5a cells respond to multiple sugars in addition to trehalose (29). However, it is also unclear whether other sugar-activated Grs are activated by a single ligand as appears to be the case for Gr5a.

Here we used a rapid screening approach to identify other Grs that may be coexpressed with Gr5a. We found that the RNAs encoding all seven Grs, which were most related phylogenetically to *Gr5a* (*Gr64a-f* and *Gr61a*), were coexpressed with the *Gr5a* GRNs. Because these Gr5a-expressing GRNs function in the response to sweet but not bitter tastants, those receptors that are coexpressed with Gr5a would be candidate sugar receptors. We generated *Gr64a*-deficient flies and showed that this receptor is required for the attractive behavioral responses to glucose, which is the most widely distributed sugar in plants and animals, as well as to the disaccharides, sucrose, and maltose. We also showed that flies missing *Gr64a* do not generate action potentials in response to these sugars. These results indicate that multiple Grs are coexpressed with Gr5a and that the one receptor/one ligand paradigm for Gr5a does not apply to all sugar receptors in *Drosophila*.

Results

mRNA Tagging Approach for Identification of Candidate Sugar Receptors. To identify Grs that were candidate sugar receptors, we tested whether any Gr RNAs were coexpressed with *Gr5a*. Given that *in situ* hybridizations to Gr RNAs have been problematic for most Gr genes (4, 5), we used an mRNA tagging approach (30) in combination with the *GAL4/UAS* system (31). The mRNA tagging approach entails purification of RNAs from specific cell populations using a FLAG-tagged poly(A)-binding protein (PABP). To screen for Grs expressed in Gr5a GRNs, we prepared extracts from dissected labella expressing *Gr5a-GAL4;UAS-PABP* or *Gr66a-GAL4;UAS-PABP* transgenes, then

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Abbreviations: Gr, gustatory receptor; GRN, Gr neuron; PABP, poly(A)-binding protein; PI, preference index.

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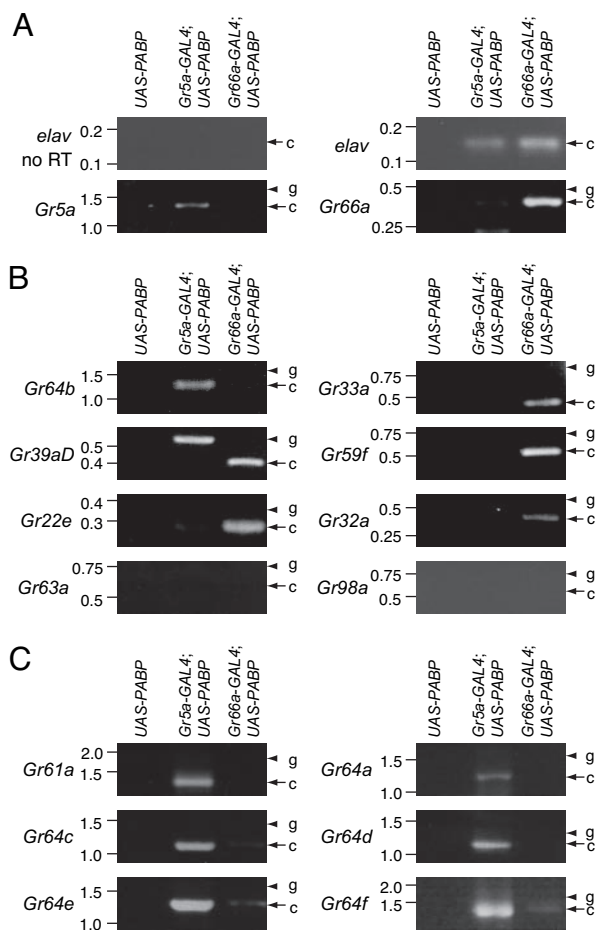


Fig. 1. RT-PCR screen for *Gr* RNAs expressed in *Gr5a*-positive GRNs. (A) Examination of *Gr5a* and *Gr66a* RT-PCR products in flies expressing *UAS-PABP* under control of the *Gr5a-GAL4* or *Gr66a-GAL4*. The RNAs were prepared from the labella of flies containing the *UAS-PABP* transgene only and from *Gr5a-GAL4;UAS-PABP* and *Gr66a-GAL4;UAS-PABP* flies. *elav* was used as an internal control. The arrows and arrowheads indicate the products generated from the reverse-transcribed mRNA and from the genomic DNA templates, respectively. The sizes of DNA markers (kb) are indicated to the left. (B) RT-PCR products for *Gr64b*, *Gr33a*, *Gr39aD*, *Gr59f*, *Gr22e*, *Gr32a*, *Gr63a*, and *Gr98a*. No RT-PCR products for *Gr63a* or *Gr98a* were detected. (C) RT-PCR products for the *Gr-S* group (*Gr64* cluster genes and *Gr61a*).

immunoprecipitated the FLAG-PABP with anti-FLAG antibodies and isolated the mRNAs that were pulled down with the PABP (see *Methods*).

To test the efficacy of the mRNA tagging, we performed RT-PCR using primers specific for *Gr5a* and *Gr66a*. As a control, we performed RT-PCR using primers specific for a panneuronally expressed gene, *elav* (32), which was therefore detected in both *Gr5a*- and *Gr66a*-expressing GRNs (Fig. 1A). In contrast to these results, the *Gr5a* signal was much higher using RNA prepared from *Gr5a-GAL4;UAS-PABP* than from *Gr66a-GAL4;UAS-PABP* flies (Fig. 1A). Conversely, the *Gr66a* product was found primarily by using RNA from *Gr66a-GAL4;UAS-PABP* flies (Fig. 1A). These results indicated that the mRNA tagging approach may be an effective assay to examine whether other *Gr*s are expressed in either *Gr5a*- or *Gr66a*-expressing GRNs.

Initially we surveyed the expression of a set of eight *Gr*s that were distributed among a variety of branches within the *Gr* family tree [supporting information (SI) Fig. 5] (22). Two of the *Gr* RNAs (*Gr22e* and *Gr32a*) were predicted to be enriched in *Gr66a*-positive

neurons because the corresponding *GAL4* reporters have been shown to be expressed in subsets of *Gr66a* GRNs (4, 5). In addition, we examined *Gr63a* expression, which was unlikely to be coexpressed with either *Gr5a* or *Gr66a* because this *Gr* encodes a CO₂ receptor (33, 34). *Gr64b* belonged to a distinct branch that included the seven *Gr*s most related to *Gr5a* (28–45% amino acid identities; referred to here as the *Gr-S* group).

We found that *Gr64b* was the only one among the eight surveyed that was enriched in *Gr5a-GAL4;UAS-PABP* flies (Fig. 1B). In contrast, five of the *Gr* RNAs were found primarily in the RNA prepared from *Gr66a-GAL4;UAS-PABP* (Fig. 1B), including three whose expression had not been previously characterized (*Gr33a*, *Gr39aD*, and *Gr59f*). No *Gr63a* product was detected in either RNA sample (Fig. 1B), which was expected because *Gr63a* is a CO₂ receptor. A *Gr98a* RT-PCR band also was not detected, which might be because of its low expression level.

Because *Gr64b* was among the group of seven *Gr*s most related to *Gr5a*, we tested whether the remaining six members of the *Gr-S* group were enriched in *Gr5a-GAL4;UAS-PABP* flies. We found that the RT-PCR products of all *Gr-S* RNAs (*Gr61a* and *Gr64a-f*) were expressed predominately in *Gr5a-GAL4;UAS-PABP* flies (Fig. 1C). These included *Gr64a* and *Gr64e*, despite the report that *Gr64a*- and *Gr64e*-*GAL4* reporter expression was not detected in the labellum (4). Thus, of the 14 *Gr*s tested, all seven *Gr-S* RNAs but none of the other *Gr*s were enriched in *Gr5a-GAL4;UAS-PABP* flies.

***Gr64a* Is Required for the Behavioral Responses to Multiple Sugars.**

We queried the available P-element collections for insertions that disrupted *Gr61a* or one of the *Gr64a-f* genes. One P-element was available (GE24923; Genexel, Daejeon, South Korea), which inserted 336 bp upstream from the 5' end of *Gr64a* (Fig. 2A). Flies homozygous for the GE24923 insertion were viable and fertile and showed normal responses to all sugar tested and normal levels of the *Gr64a* RT-PCR product (data not shown). The flies also showed no significant differences from wild-type flies (Canton S) in their sugar preferences (Fig. 3A and SI Fig. 6).

To generate a deletion flanking the 3' end of GE24923, we genetically introduced the transposase because mobilization of P-elements can result in deletions flanking the site of the original insertion. After screening genomic DNA from ≈ 800 lines by PCR, we identified one with a 3.1-kb deletion, which removed all of *Gr64a* and 868 bp at the 5' end of *Gr64b*. In addition, a 2.0-kb portion of the flanking gene, *CG11594*, was duplicated and inserted in the deleted region (Fig. 2A). The original *CG11594* gene was not disrupted by the deletion (Fig. 2B). This mutant line is referred to as *Gr64^{ab}*.

We examined whether the *Gr64* cluster genes were expressed in *Gr64^{ab}* homozygous flies by performing RT-PCR using total RNA prepared from labella. We found that expression of the *Gr64a* and *Gr64b* mRNAs was eliminated, whereas the mRNAs from all of the other four genes (*Gr64c*, *Gr64d*, *Gr64e*, and *Gr64f*) in the cluster were still produced (Fig. 2C).

To determine whether *Gr64^{ab}* flies had a defect in detecting sugars, we used a two-sugar choice test. When presented with a choice between 5 mM sucrose and 2 mM fructose, wild-type flies displayed a very strong preference to consume the sucrose (Fig. 3C) [preference index (PI) = 0.97 ± 0.03 ; $n = 6$]. This bias for selecting sucrose was not due to an inability to detect fructose, because wild-type flies selected the higher of two concentrations of fructose (5 mM versus 2 mM) (Fig. 3A) (PI = 0.88 ± 0.04 ; $n = 3$), similar to the situation when offered two different concentrations of sucrose (5 mM versus 1 mM) (Fig. 3A) (PI = 0.92 ± 0.03 ; $n = 5$).

In contrast, *Gr64^{ab}* flies preferred fructose to sucrose (Fig. 3C) (PI = 0.10 ± 0.05 ; $n = 6$). This suggested that the response to sucrose was reduced or eliminated. In addition, *Gr64^{ab}* flies failed to show a bias in favor of the 5 mM over the 1 mM sucrose

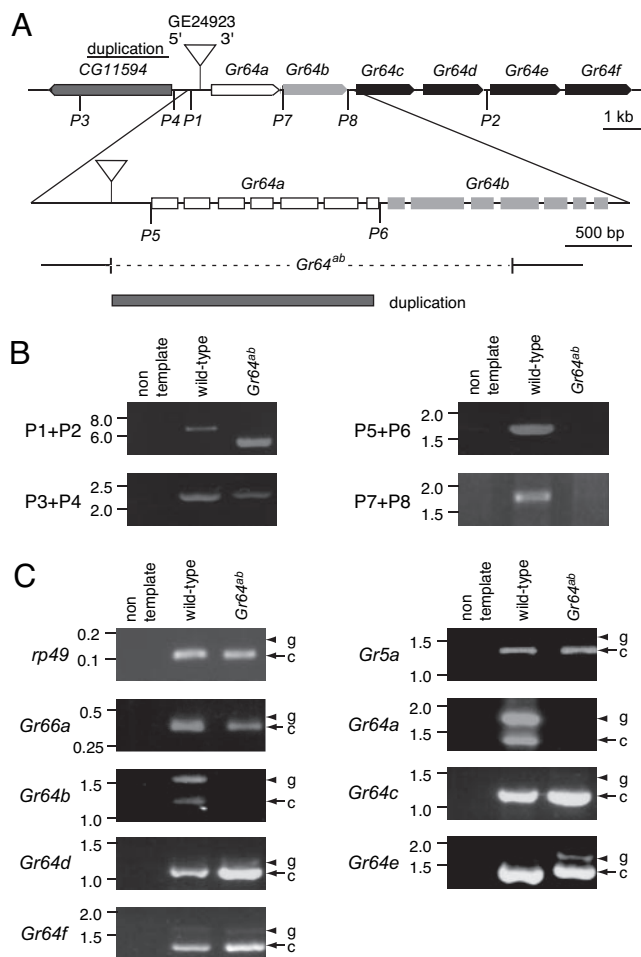


Fig. 2. Generation of the *Gr64^{ab}* mutant. (A) The genomic region encoding *Gr64a-f*. The top horizontal line indicates the portion of the 64A4 region that includes *Gr64a-f* and *CG11594*. The pointed end indicates the 3' end of each gene. The inverted triangle indicates the P-element (GE24923) insertion site. P1 and P2 indicate the locations of the PCR primers used in the primary screen for a deletion flanking GE24923. P3–P8 indicate the primers used in the subsequent analyses. The introns/exons of *Gr64a* and *Gr64b* are shown at a higher resolution below. One line (*Gr64^{ab}*) contained a 3.1-kb deletion, which extended from the P-element insertion site to the fourth exon of *Gr64b* (dashed line), and a duplication of a 2.0-kb portion of *CG11594* as indicated. (B) PCR analyses of the *Gr64^{ab}* region. The PCR product using the P1 and P2 primers generated 6.2- and 5.0-kb products from wild type and *Gr64^{ab}*, respectively. The P3 and P4 primers, which flank the duplicated region, produce the same products in wild type and *Gr64^{ab}*. The P5/P6 and P7/P8 primer pairs produced products only in wild type because of the deletion in *Gr64^{ab}*. The precise alterations were determined subsequently by DNA sequencing. (C) Analyses of *Gr64a-f* RT-PCR products. The control RT-PCR products (*rp49*, *Gr5a*, and *Gr66a*) were not affected in *Gr64^{ab}*. The arrows and arrowheads indicate the products generated from the reverse-transcribed mRNA and from the genomic DNA templates, respectively. DNA size markers (kb) are shown.

(Fig. 3A) ($PI = 0.45 \pm 0.06$; $n = 5$) but retained the ability to select the 5 mM over 2 mM fructose (Fig. 3A) ($PI = 0.86 \pm 0.02$; $n = 3$). Thus, the behavioral response to sucrose, but not fructose, was defective in *Gr64^{ab}* flies.

We used the two-sugar choice assay to test the responses of *Gr64^{ab}* flies to other sugars. In the case of wild-type flies, there was a strong propensity to consume 20 mM glucose, 10 mM maltose, 20 mM trehalose, or 20 mM arabinose over 2 mM fructose (Fig. 3C) ($PI = 0.98 \pm 0.01$, 0.94 ± 0.04 , 0.89 ± 0.03 , and 0.89 ± 0.01 , respectively; $n = 3-6$). The *Gr64^{ab}* flies also exhibited a normal

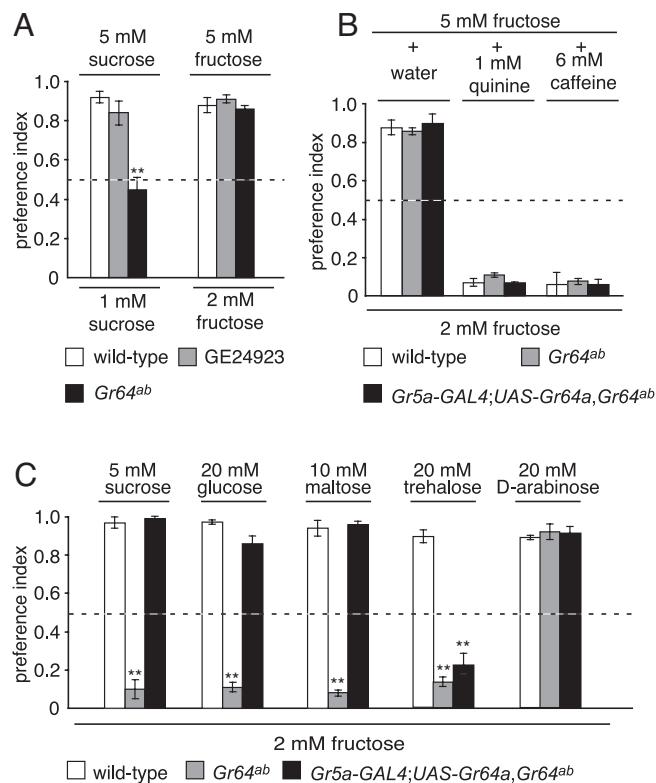


Fig. 3. Two-way choice tests showing that *Gr64a* was required for the behavioral responses to multiple sugars. PI values of 1.0 or 0 indicate complete preferences for one or the other tastant, and a PI of 0.5 indicates a lack of preference. (A) Flies were allowed to choose between two concentrations of the same sugar (sucrose or fructose). (B) *Gr64a* was not required for the avoidance of quinine or caffeine. The flies were given the choice between 2 mM fructose or 5 mM fructose plus either 1 mM quinine or 6 mM caffeine. (C) Two-sugar competition assay. The following fly stocks were allowed to choose between 2 mM fructose and higher concentrations of other sugars as indicated: (i) wild type (Canton S), (ii) *Gr64^{ab}*, and (iii) *Gr64^{ab}* flies containing the *Gr5a-GAL4;UAS-Gr64a* transgenes. The dashed line indicates a PI of 0.5. Statistically significant differences from wild type were checked by using the unpaired Student *t* test (**, $P < 0.01$).

preference for 20 mM arabinose over 2 mM fructose (Fig. 3C) ($PI = 0.92 \pm 0.04$; $n = 3$). However, unlike wild type, the *Gr64^{ab}* mutant preferred fructose over either glucose or maltose (Fig. 3C) ($PI = 0.11 \pm 0.03$ and 0.08 ± 0.02 , respectively; $n = 6$). Also, in contrast to wild type, *Gr64^{ab}* selected fructose rather than trehalose (Fig. 3C) ($PI = 0.15 \pm 0.02$; $n = 6$), which was surprising given that trehalose is the ligand that activates *Gr5a* (26–28).

Bitter compounds, such as quinine and caffeine, are aversive to wild-type flies, and this response is mediated by GRNs that express *Gr66a* but not *Gr5a* (4, 5). Consistent with the expression of the *Gr64a* and *Gr64b* RNAs in *Gr5a*-expressing GRNs, we found that *Gr64^{ab}* flies showed the normal aversion to quinine and caffeine (Fig. 3B) (wild type, $PI = 0.07 \pm 0.02$ and 0.06 ± 0.03 , respectively; *Gr64^{ab}*, $PI = 0.11 \pm 0.01$ and 0.08 ± 0.02 ; $n = 3$, respectively).

Because both *Gr64a* and *Gr64b* were deleted in the *Gr64^{ab}* mutant, we addressed whether the defects in the behavioral responses to sucrose, glucose, maltose, and trehalose detection were due to loss of *Gr64a* or *Gr64b*. We generated *UAS-Gr64a* and *UAS-Gr64b* transgenes and expressed them in *Gr64^{ab}* flies under the control of the *Gr5a-GAL4*. The *UAS-Gr64b* and *Gr5a-GAL4* transgenes did not restore a normal response to any sugar, including trehalose (data not shown). However, introduction of the *UAS-Gr64a* transgene, in combination with the *Gr5a-GAL4*, fully restored the preferences for sucrose, glucose,

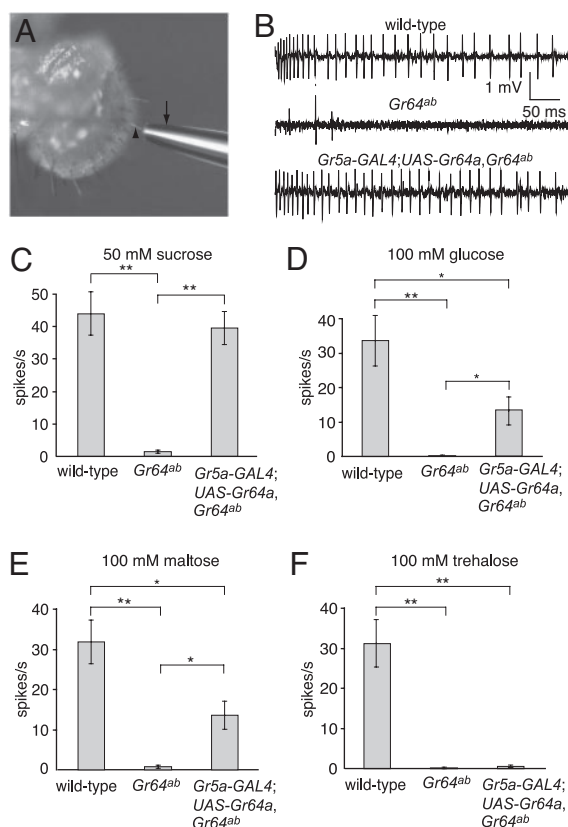


Fig. 4. *Gr64a* was required for sucrose-, glucose-, and maltose-induced action potentials. Tip recordings were performed on L4 sensilla of the indicated genotypes. (A) Recording electrode (arrow) placed over an L type bristle (arrowhead). (B) Sample tip recordings using 50 mM sucrose. (C–F) Average frequencies of action potentials (spikes per second) in response to the indicated sugars. The averages were based on data collected between 50 msec and 1,050 msec after application of the sugars ($n = 5–7$). Statistically significant differences between the indicated pairs of data were checked by using the unpaired Student *t* test (*, $P < 0.05$; **, $P < 0.01$).

or maltose over fructose (Fig. 3C) ($PI = 0.99 \pm 0.01$, 0.86 ± 0.04 , and 0.96 ± 0.02 , respectively; $n = 6$) but not the trehalose response ($PI = 0.22 \pm 0.06$; $n = 6$). Introduction of both *UAS-Gr64a* and *UAS-Gr64b* with *Gr5a-GAL4* also did not rescue the trehalose response (data not shown). Nevertheless, the trehalose defect appeared to be associated with the *Gr64^{ab}* mutation, rather than a background mutation, because the phenotype was observed in flies containing the *Gr64^{ab}* chromosome *in trans* with either of two deficiency chromosomes that span the *Gr64* locus (SI Fig. 7). Furthermore, introduction of a transgene that included the entire *CG11594* genomic region did not rescue any aspect of the *Gr64^{ab}* phenotype (data not shown). These results demonstrate that *Gr64a* functions in the detection of sucrose, glucose, and maltose in *Gr5a*-expressing GRNs. As with many Grs, we were unable to detect expression of *Gr64a* RNA by *in situ* hybridizations, presumably because of low expression levels.

Sucrose-, Glucose-, and Maltose-Induced Action Potentials Require *Gr64a*.

To address whether the electrophysiological response to sugars was defective in *Gr64a*-deficient flies, we assayed tastant-induced action potentials in the GRNs by performing tip recordings (Fig. 4A). In *Drosophila*, sugar-induced action potentials can be detected in the GRNs in L, I, or S type bristles, although the highest frequencies and rates of response occur in L type sensilla (35). Therefore, we applied sugars to L type

bristles and compared the frequencies of action potentials in wild type and in the *Gr64^{ab}* mutant. In wild-type flies, sucrose, glucose, maltose, and trehalose stimulated high frequencies of action potentials (Fig. 4B–F) (44.0 ± 6.7 , 34.4 ± 8.7 , 31.8 ± 5.5 , and 33.4 ± 5.7 , respectively; $n = 5–7$), whereas the frequencies of action potentials induced by these sugars in *Gr64^{ab}* were dramatically reduced (Fig. 4B–F) (1.4 ± 0.5 , 0.2 ± 0.2 , 0.8 ± 0.3 , and 0.8 ± 0.5 , respectively; $n = 5–7$). Consistent with the behavioral assays, the reduction in action potentials in response to trehalose was not rescued by either of the wild-type *Gr64a* or *Gr64b* transgenes (Fig. 4F) (0.6 ± 0.04 ; $n = 5$ and data not shown). In contrast, introduction of the *Gr5a-GAL4* and *UAS-Gr64a* transgenes restored action potentials to various extents in response to sucrose, glucose, and maltose (Fig. 4B–E) (39.6 ± 5.0 , 12.8 ± 4.9 , and 13.6 ± 3.4 , respectively; $n = 5–7$). These results indicated that *Gr64a* was required in GRNs for the detection of sucrose, glucose, and maltose.

Discussion

Before the current analysis, the only known *Drosophila* Grs linked to specific tastants were *Gr5a* (26–28) and *Gr66a* (25), which are required for the responses to trehalose and caffeine, respectively. All nine of the other Grs shown to be expressed in the labellum appeared to be coexpressed in subsets of *Gr66a*-expressing GRNs (4, 5). Thus, there were many candidate bitter Grs. However, the identities of candidate sugar-responsive Grs, or whether any Grs were coexpressed with the trehalose receptor, were not known. An additional question is whether other sugar receptors were tuned to single or multiple sugars.

Because in most cases detection of *Gr* RNAs by *in situ* hybridizations has been unsuccessful, we screened for Grs that are coexpressed with *Gr5a* using an mRNA tagging approach. We found that expression of the seven Grs that were most related to *Gr5a* (*Gr64a-f* and *Gr61a*) were enriched in *Gr5a*-expressing GRNs. Thus, all eight members of this phylogenetically distinct group, referred to as Gr-S, are candidate sugar receptors. None of the other 14 Grs analyzed here or in previous studies are coexpressed with *Gr5a* (4, 5). Because T1R2 and T1R3 seem to account for all of the sugar responses in mammals (16, 36), it may be that there is a larger repertoire of sugar receptors in *Drosophila*.

To test the proposal that Gr-S receptors other than *Gr5a* are sugar receptors, we examined the requirement for *Gr64a*. An additional question is whether other *Drosophila* sugar receptors are activated by one or multiple sugars. In contrast to the mammalian sugar receptors, *Gr5a* was reported to be specifically activated by trehalose (26–28). This observation, in combination with the larger number of candidate sugar receptors in flies than mammals, raised the possibility that each Gr-S member may respond primarily to one sugar. However, we found that the *Gr64a* gene was required *in vivo* for the responses to the monosaccharide, glucose, and the disaccharides, sucrose and maltose, each of which includes at least one glucose subunit. *Gr64a* was not essential for the responses to all sugars, because the *Gr64^{ab}* flies responded normally to the monosaccharides, fructose and arabinose. Although the defects in the behavioral responses to sucrose, glucose, and maltose were completely rescued by expression of a wild-type *Gr64a* transgene, only the sucrose response was fully rescued as assayed by tip recordings. The rescues of the electrophysiological responses to glucose and maltose were significant but did not restore the same frequencies of action potentials as in wild type. Although the explanation for this result is unclear, similar findings were reported for rescue of the trehalose deficits in the *Gr5a^{Δ5}* allele by a wild-type transgene (28). Whereas the behavioral phenotype in *Gr5a^{Δ5}* was restored entirely, the rescue of the electrophysiological response to trehalose was partial.

The trehalose response was also greatly reduced in *Gr64^{ab}* mutant flies. This result was unexpected because the response to

this sugar is nearly eliminated in the *Gr5a* mutant, and *Gr5a* is sufficient to confer trehalose sensitivity in S2 cells (26). Nevertheless, there is small residual trehalose response in the *Gr5a* mutant (28). The trehalose defect in *Gr64^{ab}* flies did not appear to arise from a background mutation in *Gr5a*, because the phenotype was observed in flies in which the *Gr64^{ab}* deletion was placed *in trans* with deficiencies that spanned the *Gr64* locus. Thus, the question arises as to the identity of the second trehalose receptor. The deletion in *Gr64^{ab}* disrupts both *Gr64a* and *Gr64b*, and introduction of a wild-type *UAS-Gr64a* transgene under the control of the *Gr5a-GAL4* restores normal responses to sucrose, glucose, and maltose, but not to trehalose. Thus, *Gr64b* may be a trehalose receptor. However, a *UAS-Gr64b* transgene alone or in combination with the *UAS-Gr64a* failed to restore a trehalose response in flies containing the *Gr5a-GAL4* (Y.J. and C.M., unpublished data), suggesting that either *Gr64b* is not a trehalose receptor or the transgene is nonfunctional.

A general issue concerning the *Drosophila* Grs is whether they typically form homo- or heteromultimers. An indication that at least some Grs form obligatory heteromultimers is that misexpression of just one of the two CO₂ receptor genes, *Gr21a* or *Gr63a*, in CO₂-insensitive antennal neurons is insufficient to confer CO₂ sensitivity to these cells. However, coexpression of both *Gr21a* and *Gr63a* induces CO₂ responsiveness (33, 34). Misexpression of just *Gr64a* in *Gr66a* GRNs did not elicit an aversive response to sucrose, glucose, or maltose or result in sugar-induced action potentials in *Gr66a* GRNs (Y.J. and C.M., unpublished data). Similarly, expression of the caffeine receptor *Gr66a* in *Gr5a*-expressing cells does not confer caffeine sensitivity to these cells (S.J.M. and C.M., unpublished data). These results raise the possibility that these and possibly other taste receptors in flies are obligatory heterodimers, as is the case for the CO₂ receptors. The other Gr-S members would appear to be the best candidates for forming heteromultimers with *Gr64a*. Finally, we propose that the mRNA tagging approach applied here can be extended to identify pairs of Grs that are expressed together in smaller subsets of GRNs and would therefore be excellent candidates for forming heteromultimers.

Methods

Genetics, Fly Stocks, and Constructs. The GE24923 P-element (Genexel), which inserted 366 bp 5' of *Gr64a*, was mobilized by genetically introducing the transposase using the $\Delta 2-3$ line (37): *w;Sp/CyO; ry Sb¹ P{ry[+t7.2] = $\Delta 2-3$ }99B/TM6B* (Bloomington *Drosophila* Stock Center, Bloomington, IN). To identify the *Gr64^{ab}* deletion, we screened ≈ 800 lines by PCR as previously described (38) using the following primers: P1, 5'-TTATTA-GAAGCGCGCACACCTA CTC-3'; P2, 5'-ACAAGGATATC-CAGCGAAAGCGCA-3' (Fig. 2A).

To create the P[*UAS-Gr64a*] and P[*UAS-Gr64b*] transgenes, we amplified the coding regions of *Gr64a* and *Gr64b* from total labellar RNA by RT-PCR and subcloned the cDNAs into the pUAST vector (31). The clones were sequenced, confirming that no mutations were introduced. The transgenic lines were generated by using standard procedures, and the transgenes were crossed into the *w¹¹¹⁸;Gr64^{ab}* background. The P[*Gr5a-GAL4*] and P[*Gr66a-GAL4*] transgenic flies were kindly provided by H. Amrein (Duke University Medical Center, Durham, NC).

Chemicals. Sucrose, glucose, maltose, D-arabinose, quinine, caffeine, and tricholine citrate were purchased from Sigma-Aldrich (St. Louis, MO), and trehalose was from Fluka (St. Gallen, Switzerland).

mRNA Tagging. The mRNA tagging protocol was modified from that described previously (30). Approximately 400 fly labella were dissected and fixed in 1 ml of PBS with 1% formaldehyde and 0.5%

Nonidet P-40 for 30 min at 4°C. A total of 140 μ l of 2 M glycine was added, and the samples were incubated for an additional 5 min at 4°C. The samples were washed three times with 1 ml of PBS and homogenized in 0.8 ml of homogenization buffer (HB: 150 mM NaCl/50 mM Hepes, pH 7.6/1 mM EGTA/15 mM EDTA/10% glycerol). Immediately before addition of the HB, we added the following to the HB at the indicated final concentrations: 8 mM vanadyl ribonucleoside complex (Sigma-Aldrich), 50 units/ml SUPERase-In (Ambion, Austin, TX), and a protease inhibitor mixture tablet (one tablet per 50 ml; Roche, Indianapolis, IN). The homogenate was then sonicated for 1 min at 30% intensity (using a Fisher Sonic Dismembrator, Model 500) and cleared by centrifugation for 10 min at 13,000 $\times g$. Anti-FLAG-M2 affinity agarose beads (Sigma-Aldrich) were washed four times with HB at 4°C by centrifugation for 1 min at 1,500 $\times g$. To coimmunoprecipitate the FLAG-tagged PABP and the associated polyadenylated mRNAs, we added 100 μ l of anti-FLAG-M2 affinity agarose beads to the cleared homogenate, which was then incubated for 2 h at 4°C. The beads were washed four times with the HB at 4°C and incubated in 100 μ l of elution buffer (50 mM Tris-HCl, pH 7.0/10 mM EDTA/1.3% SDS/50 units/ml SUPERase-In) at 65°C for 30 min to reverse the RNA::PABP crosslink. A total of 100 μ l of eluant was treated with 400 μ l of TRIzol (Invitrogen, Carlsbad, CA) and 100 μ l of chloroform, and the RNA extracted was finally dissolved in 40 μ l of RNase-free water. The P[*UAS-hPABP-FLAG*] transgenic flies used for these experiments were kindly provided by R. L. Davis (Baylor College of Medicine, Houston, TX).

RT-PCR Amplification of *Gr* RNAs. Two types of RNA preparations were used for performing the RT-PCR: (i) the mRNA tagging approach was used to prepare the RNA as described above (Fig. 1), or (ii) 100 labella were dissected and total RNA was extracted by using TRIzol reagent (Invitrogen) (Fig. 2). The RNA samples were treated with DNaseI (Invitrogen) before the RT-PCRs were performed. Each pair of primers to a given *Gr* was designed to span at least one intron so that the products derived from mRNA and genomic DNA could be discriminated on the basis of size. Because *elav* has no introns, we used a no-reverse-transcriptase control to confirm that the RT-PCR product was derived from mRNA instead of genomic DNA. A OneStep RT-PCR Kit (catalog no. 210212; Qiagen, Valencia, CA) was used for the RT-PCR. To perform the RT-PCR with total RNA we amplified for 30–35 cycles, and for the mRNA tagging experiments we used 40 cycles.

Behavioral Assays. The two-way choice assays were performed by using modifications of previously described procedures (25, 39). For each assay, 30 flies (3–6 days old) were starved overnight on 1% agarose and then introduced into 72-well microtiter dishes filled with two types of test mixtures placed in alternating wells. Each test mixture contained 1% agarose and either blue dye (0.125 mg/ml brilliant blue FCF, catalog no. 027-12842; Wako Chemical, Richmond, VA) or red dye (0.2 mg/ml sulforhodamine B, catalog no. S9012; Sigma-Aldrich) and a test tastant. For the attractive tests, the wells were filled with (i) one sugar at two different concentrations or (ii) 2 mM fructose and a different sugar at a higher concentration (5–20 mM). A higher concentration of the second sugar was selected if it induced a PI of >0.8 . The avoidance assays were conducted by using 2 mM fructose versus 5 mM fructose plus either 1 mM quinine or 6 mM caffeine. After allowing the flies to feed for 90 min at room temperature in the dark, the animals were frozen in the dishes at -20°C , and the numbers of flies with blue (N^{B}), red (N^{R}), or purple (N^{P}) abdomens were assessed by visual inspection. In those cases in which the colors were difficult to judge, the guts were dissected. If the amount of red dye was between 50% and 150% of the blue dye, the color was scored as purple. If the red dye was $>150\%$ or $<50\%$ of the blue dye, the fly was counted

as red and blue, respectively. All behavioral assays (30 flies per test) were performed three to six times. The PI values were calculated according to the following equation: $PI = (N^B + 0.5 N^P)/N^{Total}$ or $(N^R + 0.5 N^P)/N^{Total}$. The dyes did not cause preference changes, because no differences in PIs were caused by switching the dyes. For example, in those assays in which wild-type flies were allowed to choose between 5 mM sucrose and 2 mM fructose, the PI values were 0.96 ± 0.04 when blue and red dyes were used for sucrose and fructose, respectively, and 0.97 ± 0.03 when blue and red dyes were used for fructose and sucrose, respectively. The wild-type control was Canton S, which showed behavioral responses indistinguishable from the parental P-element insertion line (GE24923) (SI Fig. 6).

Electrophysiology. Tip recordings on labellar bristles were performed according to procedures similar to those previously described (25). Briefly, to provide a reference electrode and to stabilize the fly for the recordings, we inserted a glass capillary with Ringer's solution into the abdomen so that it extended through to the fly head. The electrolyte used in the recording electrode (10–20 μ m in diameter) was 30 mM tricholine citrate.

The recordings were performed on the L4 sensilla (flies ≤ 1 day after eclosion). The signals were collected and amplified from the recording electrode through a preamplifier (TastePROBE; Syntech, Hilversum, The Netherlands) and a 100- to 3,000-Hz band-pass filter. Autospike 3.1 software (Syntech) was used to acquire the action potentials (9.6-kHz sampling rate) and to analyze the frequencies. For most sugars we used 100 mM, which is the concentration typically used in previous analyses (28, 35). We used 50 mM sucrose because this sugar induced a much higher frequency of action potentials than other sugars. All recordings using a given genotype and tastant were performed five to seven times.

Data Analyses. All error bars represent SEMs. Unpaired Student's *t* tests were used to check for significant differences between the indicated pairs of data (*, $P < 0.05$; **, $P < 0.01$).

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