Physiological and Behavioral Responses to Optogenetic Stimulation of PKD2L1+ Type III Taste Cells

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Abstract

Type III taste cells in mammalian taste buds are implicated in the detection and communication of sour and some salty stimuli, as well as carbonation and water. With this variety of proposed roles, it is unclear what information activated type III cells are communicating to the CNS. To better elucidate the role of type III cells in the taste bud, we use a type III cell-specific protein (polycystic kidney disease 2-like 1) to drive Cre-dependent expression of light-sensitive channelrhodopsin (Ai32) in mouse type III taste cells. Activation of these cells with light produces a taste nerve response in both the chorda tympani and glossopharyngeal nerves, and elicits a slight but significant aversion in two-bottle preference tests in both male and female mice. Unlike previous reports (Zocchi et al., 2017), our mice did not react to blue light stimulation with sustained drinking responses. These data suggest that type III cells are capable of communicating the presence of aversive stimuli in the oral cavity, which is in line with their responsiveness to sour and high concentrations of salt stimuli.

Key words: optogenetics; taste; type III taste cells

Significance Statement

The sour-sensing type III cells of the mammalian taste bud are historically difficult to study in vivo, as sour stimulation of the tongue causes intracellular acidification in all epithelial cells, not just type III cells. The present study circumvents this issue by using Cre-dependent expression of channelrhodopsin in type III cells, so that they can be activated by blue light, rather than acid (sour). Our data suggest that type III cells communicate an aversive signal to the CNS.

Introduction

Taste buds are the sensory end organs of the gustatory system, allowing organisms to discern nutritious energy sources from potentially dangerous ones. In most mammals, these clusters of sensory cells detect at least five basic taste modalities: bitter, sweet, umami, sour, and salty. Some sensory cells in the taste bud process discreet modalities; each type II cell in the taste bud transduces only bitter, sweet, or umami stimuli, and com-
municates this information to afferent nerve fibers via nonvesicular ATP release (Finger et al., 2005; Taruno et al., 2013; Romanov et al., 2018). Type I cells are thought to play primarily a support role in the taste bud similar to glial cells, with enzymes for uptake and degradation of transmitters (Lawton et al., 2000; Bartel et al., 2006; Dvoranyanchikov et al., 2009). Here we focus on type III cells, which are implicated in a variety of functions in the taste bud.

Type III cells are elongated, spindle-shaped cells featuring synaptic connections with afferent nerve fibers (Murray et al., 1969). Since their identification as distinct ultrastructural components of the taste bud, these cells have been associated with several physiologic properties. Type III cells express the polycystic kidney disease 2-like 1 (PKD2L1) ion channel (Kataoka et al., 2008), and, although the function of this protein remains unclear, it has been used as a marker of type III cells and the Pkd2l1 gene has been used as a Cre driver to manipulate gene expression. Genetic deletion of PKD2L1-expressing taste cells eliminates chorda tympani nerve responses to sour (acidic) taste stimuli (Huang et al., 2006), but type III cells have also been implicated in responses to carbonation (Chandrashekar et al., 2009) and high concentrations of salt (Oka et al., 2013; Lewandowski et al., 2016), all of which are considered aversive modalities. However, a recent study (Zocchi et al., 2017) entailing expression of channelrhodopsin-2 (ChR2) challenged the negative valence of type III cells, suggesting instead that type III cells primarily mediate water detection and drive drinking behavior. The mouse used in their study was produced from a BAC-transgene containing the Pkd2l1 locus, which was then used to drive ChR2 in type III cells. On blue light stimulation of the tongue, mice exhibited continuous licking, even in the absence of water in the sipper tube. Zocchi et al. (2017) suggested that the averseness of acids may result not from the activation of type III cells, but from additional mechanisms of acid detection in the tongue, such as trigeminal afferents.

We have developed a similar mouse to manipulate gene expression in type III cells, also using the Pkd2l1 gene as a Cre driver. However, we made our mice by knockin of an IRES Cre recombinase construct directly following the stop codon (Ye et al., 2016). This mouse was characterized and used to knock down the potassium channel K\textsubscript{p}2.1, validating its role in sour taste transduction (Ye et al., 2016). In the present study, we have used this mouse to re-examine the role of type III cells in taste behavior. We crossed our Pkd2l1-Cre mice to floxed ChR2 mice, validated the expression of ChR2 immunohistochemically, and characterized the nerve response resulting from light stimulation of the tongue. To assess behavior, we used two-bottle preference tests and brief-access Lickometer tests during optogenetic stimulation of the tongue. Our results indicate that our Pkd2l1-Cre, ChR2 mice avoid blue light stimuli in two-bottle preference tests and, contrary to the previous study, show no sustained licking responses to light stimulation. These data suggest that selective activation of PKD2L1\textsuperscript{+} type III cells elicits primarily a negative valence, as would be expected for a sour stimulus.

Materials and Methods

Mice

All mice were housed at the University of Colorado Anschutz Medical Campus on a 12 h light/dark cycle and had continual access to standard chow. The Animal Care and Use Committee at the University of Colorado School of Medicine approved all procedures. To drive the expression of the light-sensitive ChR2 in PKD2L1\textsuperscript{+} type III cells, we crossed a Cre-dependent ChR2 mouse line (stock #012569, The Jackson Laboratory) to a Pkd2l1-Cre mouse created in house (Ye et al., 2016). This knock-in mouse features an IRES Cre-recombinase construct directly following the Pkd2l1 coding sequence. For behavioral experiments, littermate controls lacked one or both of the necessary alleles for ChR2 expression in PKD2L1\textsuperscript{+} type III cells.

Perfusion/fixation

To fix and obtain taste tissues, mice were anesthetized with sodium pentobarbital via intraperitoneal injection at 50 mg/kg and transcardially perfused with 4% paraformaldehyde (PFA; catalog #158127, Sigma-Aldrich). Tissues were extracted and immersed in 4% PFA for 1.5–5 h. Tissues were then transferred to a 20% sucrose solution overnight at 4°C before being mounted in optimal cutting temperature compound (Thermo Fisher Scientific) and cut into 12–16 μm slices via cryostat. Tissue was collected onto charged slides (Tanner Scientific) in a 1:10 series and stored at −20°C.

Immunohistochemistry

Before antibody staining, slides were washed in 0.1 M PBS (monobasic sodium phosphate, catalog #S-5011, Sigma-Aldrich; dibasic sodium phosphate, catalog #S-0876, Sigma-Aldrich; sodium chloride, catalog #S-7653, Sigma-Aldrich) three times for 10 min on a gentle shaker. A blocking solution of 2% normal donkey serum in blocking buffer for 3 h, in darkness, at room temperature (37°C) was applied at room temperature, in a 1:10 series and stored at −20°C. All mice were housed at the University of Colorado Anschutz Medical Campus on a 12 h light/dark cycle and had continual access to standard chow. The Animal Care and Use Committee at the University of Colorado School of Medicine approved all procedures. To drive the expression of the light-sensitive ChR2 in PKD2L1\textsuperscript{+} type III cells, we crossed a Cre-dependent ChR2 mouse line (stock #012569, The Jackson Laboratory) to a Pkd2l1-Cre mouse created in house (Ye et al., 2016). This knock-in mouse features an IRES Cre-recombinase construct directly following the Pkd2l1 coding sequence. For behavioral experiments, littermate controls lacked one or both of the necessary alleles for ChR2 expression in PKD2L1\textsuperscript{+} type III cells.

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Imaging and cell counting

Tissues were imaged using a 40× oil immersion numerical aperture 1.25 lens on a Leica S5 confocal microscope, with Leica LAS AF software version 2.7.3.9723.
Cells were counted by first converting each channel into a binary image via a modified Otsu method in ImageJ (version 1.49, NIH public domain), combining channels to form a composite image, and counting cells using the ImageJ plug-in cell counter. Profiles were considered to be cells of interest if, in either marker channel, the profile had both: (1) an apparent nucleus; and (2) an elongate apical process. Profiles were considered positive for a marker if any part of the profile contained fluorescence.

Nerve recording

Mice were anesthetized with urethane at 2 g/kg (catalog #U2500, Sigma-Aldrich) and stabilized spatially with a custom head holder. A tracheotomy was performed to facilitate breathing during tongue stimulation. The chorda tympani nerve was approached ventrally, cut near the tympanic bulla, and placed on a platinum-iridium wire electrode. For glossopharyngeal recordings, the glosso- pharyngeal nerve was accessed ventrally, near the trachea. A reference electrode was inserted into nearby tissue. Nerve responses were elicited by applying tastants to the anterior or posterior tongue using a pump (Mini-pump, variable flow, Thermo Fisher Scientific). Tastants included NH4Cl 100 mM (A661, Thermo Fisher Scientific), photic stimuli were delivered via a 600 nm patch cord made by the Optogenetics and Neural Engineering Core at the University of Colorado School of Medicine [Optogenetics and Neural Engineering (ONE) Core] during constant water flow with fiber-coupled LED light sources in blue and amber wavelengths (product #M470F3 and #M595F2, Thorlabs), driven by T-Cube LED Drivers (Compact T-Cube LED Driver, Thorlabs) and a 7 MHz DDS Function Generator (product #4007B, BK Precision). Stimuli were applied for 30 s, followed by 60 s of water rinse. Nerve responses were amplified (P511, Grass Instruments), integrated over a time constant of 0.5 s, and recorded using Acknowledge software (Biopac). In most cases, responses were quantified by measuring the mean of the integrated response over 30 s from the onset of the stimulus, such that both the transient and tonic portions of the nerve response were included. For experiments involving a varied duty cycle, the length of stimulation was altered to normalize the total time of light applied to the tongue. When the duty cycle is at 50%, 5 Hz light pulses result in 100-ms-long light pulses exposing the tongue to a total of 15 s of light during the 30 s stimulation. Decreasing the duty cycle reduces the total time of light exposure, so the stimulus period was increased to compensate for this, and so on. In select experiments, purinergic receptor blocker AF353 (Afferent Pharmaceuticals) was perfused over the mouse tongue for 10 min at a concentration of 1 mM between full tastant sets to block nerve responses, and in conjunction with tastants for the remainder of the experiment.

Behavior: two-bottle preference tests

Mice were water deprived for 24 h before the test period. In two-bottle preference tests, two sipper bottles, each containing water or a tastant solution, were presented to each mouse in their home cage for 15 min to alleviate the effect of side preference; each preference score was calculated from 2 test days, for which the tastant placement in the cage was reversed. For behavioral experiments involving light, optogenetic fibers were adapted (with the assistance of the ONE Core at the University of Colorado Anschutz Campus) to fit inside sipper bottles so that light was emitted from within the sipper spout. Pulsed light (5 Hz, 7 mW, 50% duty cycle) was delivered in the same manner as for nerve recordings. All statistics and graphs were performed/generated with Prism 7 (GraphPad Software), and formatted with Illustrator and Indesign (CS6, Adobe Systems).

Behavior: lickometer studies

As in two-bottle preference tests, mice were water deprived for 24 h before the test period. A laser replaced the previously described LED system (SLOC BL473T3-050FC), allowing for a higher power output (35 mW),
similar to that used previously (Zocchi et al., 2017). As in two-bottle preference tests, the optic fiber was adapted for a sipper tube that fit into a Davis Rig (DiLog Instruments), where the mice were given access to either an empty bottle without light, an empty bottle with constant blue light, or a bottle with water but without light for 10 min, and their licks were measured by InstaCal software. All statistics and graphs were performed/generated with Prism 7 (GraphPad Software) and formatted using Illustrator and Indesign (CS6, Adobe Systems).

**Experimental design and statistical analysis**

For immunohistochemical studies designed to evaluate the efficiency of Cre-driven expression in PKD2L1\(^+\) cells, taste tissues from seven mice (three female, four male) were imaged and analyzed. Tissues from two mice (one...
female, one male) were stained and imaged to determine whether Pkd2l1-Cre drives misexpression in type II taste cells (Fig. 1). Tissues from two additional mice (one female, one male) were stained and imaged to determine whether Pkd2l1-Cre drives misexpression in taste nerve fibers or cell bodies (Fig. 2). Per mouse, 4 sections of circumvallate tissue and ~12 sections of fungiform tissue were examined, and any taste buds therein were imaged. Nerve recording experiments featured a mix of female and male mice, totaling 24 mice, presented in Figure 3. Light frequency data include three mice, all female. Light power data include four mice (two female, two male). Light duty cycle data include three mice, all female. Data collected to assess the reliability of the light response include six mice (four female, two male). To compare the reliability of the light response to traditional liquid tastant responses, a two-way ANOVA was performed between light responses and pooled liquid tastant responses (Prism 7, GraphPad Software). To determine whether amber light could also elicit a nerve response, recordings of nerve responses to blue and amber light were performed and included four mice (three female, one male). For behavioral experiments presented in Figures 4 and 5, 20 age- and sex-matched mice were used to collect data (10 control mice, 10 Pkd2l1-Cre, ChR2 mice; 14 total females, 6 total males). In Figure 4, preference scores of control mice were compared with those of Pkd2l1-Cre, ChR2 mice by unpaired t tests. In Figure 5, lick counts in the first minute of the experiments were compared between conditions by unpaired t tests. In all experiments, no differences due to the sex of the animals were observed.

Results

Channelrhodopsin expression in the Pkd2l1-Cre, ChR2 mouse

The Pkd2l1-Cre mouse successfully drives ChR2 in approximately half of PKD2L1 type III cells. In circumvallate taste tissues of five mice, ~54% of PKD2L1-immunoreactive cells were also positive for ChR2-yellow fluorescent protein (YFP) fluorescence. In fungiform tissues of the same mice, 50% of PKD2L1-immunoreactive cells displayed ChR2-YFP fluorescence (Fig. 1A–C). Importantly, ChR2-YFP fluorescence does not appear in cells immunoreactive for type II cell marker PLCβ2 (Fig. 1D). ChR2-YFP fluorescence is likewise absent from P2X3-immunoreactive nerve fibers innervating circumvallate and fungiform taste buds showing ChR2-YFP fluorescence in green and taste nerve marker P2X3-immunoreactivity (IR) in magenta. A, Confocal z-stack images of circumvallate and fungiform taste buds showing ChR2-YFP fluorescence in green and taste nerve marker P2X3-IR in magenta. B, Confocal z-stack images of geniculate ganglion cells showing ChR2-YFP fluorescence in green and taste ganglion cell marker P2X3-IR in magenta. Although green fluorescence appears in the ganglion, it seems to be restricted to satellite cells and is not present in neuronal cell bodies. Scale bars, 20 μm.
Figure 3. Optogenetic stimulation of the tongue in Pkd2l1-Cre, ChR2 mice elicits a robust, tastant-like nerve response. A, Chorda tympani (CT) and glossopharyngeal (GN) nerve responses to acidic stimuli and a 470 nm light pulse stimulus in Pkd2l1-Cre, ChR2.
late and fungiform taste buds (Fig. 2A). Examination of the geniculate ganglion, which contains cell bodies of taste neurons, reveals some ChR2-YFP fluorescent signal. This signal, however, is restricted to satellite cells that do not overlap with the P2X<sub>3</sub>-immunoreactive, rounded cell bod-
ies characteristic of taste sensory neurons (Fig. 2B). Opto-
togenic stimulation of ChR2 cells, therefore, is unlikely
to directly stimulate any non-type III cells or gustatory nerve fibers.

Optogenetic stimulation of PKD2L1<sup>+</sup> cells elicits
taste nerve responses
Pulsed blue light directed toward either the anterior or
posterior tongue of an anesthetized mouse elicits a nerve
response in the chorda tympani or glossopharyngeal
nerve similar to those elicited by tastant solutions. These
responses are only elicited in mice containing both the
Pkd2l1-Cre and Cre-dependent ChR2-YFP alleles, and
are blocked by purinergic receptor blocker AF353, which
eliminates responses to all tastant solutions (Vanden-
beuch et al., 2015; Fig. 3A–C). After testing various pa-
rameters of light presentation, we determined that blue
light pulses at 5 Hz frequency, 4–7 mW power, and a
50% duty cycle (when the light is on for half of the cycle
period) elicited a maximal chorda tympani response. For
experiments testing frequency, power and duty cycle were
kept at 7 mW and 50%, respectively. For power
dose–response curves, frequency and duty cycle were set
at 5 Hz and 50%, respectively. For duty cycle experi-
ments, frequency and power were set at 5 Hz and 7 mW
(Fig. 3D–G). To test for the consistency of this response,
we conducted several experiments repeating the same
block of tastants: NH<sub>4</sub>Cl 100 mM, citric acid 10 mM,
sucrose 500 mM, and blue light pulses at 5 Hz, 7 mW, and
50% duty cycle (each presentation of four stimuli consti-
tutes a “block”). While the consistency of tantant re-
sponses for separate experiments/animals varied (Fig.
3H, left and middle) the chorda tympani response to blue
light stimulation remained approximately as reliable as
other tastants for the course of an experiment. Citric acid
responses over several blocks did not differ from aver-
aged light responses (Fig. 3H, right). Amber light (589 nm)
does not elicit a chorda tympani nerve response (Fig. 3I).

Behavioral responses to optogenetic activation of
PKD2L1<sup>+</sup> cells
To determine whether Pkd2l1-Cre, ChR2 mice prefer or
avoid optogenetic stimulation of type III cells, we devised
an experimental paradigm to test taste preference for
light, rather than traditional tastant solution preferences
(Fig. 4A). Light was always presented in conjunction with
water in the sipper tube. Both Pkd2l1-Cre, ChR2 mice,
and littermate controls avoided increasing concentrations
of citric acid, and preferred increasing concentrations of
sucrose (Fig. 4B). In two-bottle preference tests including
light, amber light was included as a control to avoid visual
behavioral preference, as amber light is within the mouse
visual range (Jacobs 1993; Neitz and Neitz 2001) but does
not elicit a nerve response. Pkd2l1-Cre and ChR2 mice,
but not littermate controls, avoided blue light in favor of
amber light. The level of avoidance is comparable to that
of low concentrations of citric acid.

Experiments performed with a similar mouse by Zocchi
et al. (2017) showed an intriguing result—blue light stim-
ulation of the tongue during behavioral experiments elic-
ted a strong licking behavior, even in the absence of
water. As these results seem contrary to our own, we
sought to better replicate those experiments. We mea-
sured single-lick events of Pkd2l1-Cre, ChR2, and litter-
mate control mice in a Davis rig, where the mice were
presented with sipper bottles without water or blue light,
without water and with light, or with water and without
blue light. A laser light source replaced the LED light
source to better mimic the power output and geometry of
light application described in Zocchi et al. (2017). While
Pkd2l1-Cre, ChR2 mice licked more in response to light
than littermate controls, the total number of licks in the
first minute was not different between the with- and
without-light conditions, and both were significantly dif-
ferent from the with-water, without-light condition (Fig. 5).
Over the full 10 min trial, Pkd2l1-Cre, ChR2 mice pre-
sented with water were eventually satiated and disconti-
uined drinking, while mice stopped licking the spout soon
after the start of the trial when water was not present (Fig.
5B, right).

Discussion
In the present study we show that optogenetic stimu-
lation of PKD2L1<sup>+</sup> type III taste cells elicits a nerve re-

mice. B, CT nerve responses to citric acid and blue light in a Pkd2l1-Cre, ChR2 mouse before and after the application of purinergic receptor blocker AF353 to the tongue. D, CT nerve responses to a control tastant stimulus (NH<sub>4</sub>Cl) and blue light pulses at increasing frequencies (power at 7 mW, duty cycle at 50%). E, CT nerve responses to NH<sub>4</sub>Cl and blue light pulses at increasing light power (frequency at 5 Hz, duty cycle at 50%). F, CT nerve responses to NH<sub>4</sub>Cl and blue light pulses at increasing light-on duty cycle percentages (frequency at 5 Hz, power at 7 mW). G, All quantified data in frequency (n = 3), power (n = 4), and duty cycle experiments (n = 3). H, Two example experiments (left and middle) showing the consistency of the blue light response, compared with repeated blocks of 100 mM NH<sub>4</sub>Cl (black), 10 mM citric acid (CA; red), 500 mM sucrose (yellow), and blue light at 5 Hz pulses, 7 mW power, and 50% duty cycle (blue). All presentations normalized to the first presentation of that stimulus (e.g., CA responses normalized to first presentation of CA, light responses normalized to first presentation of light). Right graph shows all data (n = 6) comparing the consistency of blue light responses (blue) to CA responses (red). Blue light responses were not different from CA responses over presentation blocks (F<sub>7,110</sub> = 0.254, p = 0.9692, two-way ANOVA). I, CT nerve responses to blue (470 nm) and amber (589 nm) light. Graph on right shows all data (n = 4) of varied wavelength stimulation. Stimulus bars denote 30 s of stimulation, except for those in F, which are adjusted to normalize total time of light exposure to the tongue. All error bars denote the SEM.
response that resembles responses to tastants and is somewhat aversive to the awake, behaving mouse. Our Pkd2l1-Cre mouse drives ChR2-YFP expression in approximately half of PKD2L1-immunoreactive cells, and does not drive the expression in type II cells or taste nerve fibers or cell bodies. Thus, we are confident that this activation is specific to type III cells. Photic stimulation of the tongue elicits taste-like responses in the chorda tym-

Figure 4. Pkd2l1-Cre, ChR2 mice avoid blue light activation compared with amber control light presentation. A, Behavioral experiment sequence for (1) a typical taste stimuli two-bottle preference test, and (2) a blue v amber light two-bottle preference test. Preference score calculation at bottom. B, Taste preference curves for varying concentrations of citric acid (left) and sucrose (right) in littermate controls (gray, open), and Pkd2l1-Cre, ChR2 mice (black, filled). Scores >0.5 indicate a preference for the taste stimulus, while those <0.5 indicate an avoidance. C, Preference scores for two-bottle preference tests for 1 mM citric acid (left) and blue light (v amber, right) with littermate controls (open circles), and Pkd2l1-Cre, ChR2 mice (filled circles). Both Pkd2l1-Cre, ChR2 mice and controls slightly avoided 1 mM citric acid, but were not significantly different from each other (t = 0.31, *p = 0.7601, unpaired t test). Pkd2l1-Cre, ChR2 mice avoided blue when compared to amber light, while controls did not (t = 3.137, p = 0.0057, unpaired t test). All error bars denote 95% confidence intervals.
Pani and glossopharyngeal nerves. This blue light-induced nerve response is blocked by the purinergic receptor blocker AF353 (Vandenbeuch et al., 2015), similar to nerve responses to traditional tastant solutions. The response to blue light is maximal at a 5 Hz pulse with a 50% duty cycle at light powers of ~4 mW. Perhaps owing to the moderate expression efficiency in PKD2L1-immunoreactive cells (Fig. 1), the nerve response elicited by blue light is smaller than that elicited by 10 mM citric acid, which is near the EC50 value for citric acid (Arai et al., 2010). In two-bottle preference tests, Pkd2l1-Cre, ChR2 mice slightly avoid blue light, while control littermates are indifferent. This avoidance behavior is akin to the avoidance of low concentrations of citric acid—both are slightly, but consistently, aversive to the group of mice we tested. These data contribute to the body of evidence that identifies type III taste cells as part of a generally aversive signaling pathway (Huang et al., 2006; Oka et al., 2013).

That type III cells communicate the presence of an aversive stimulus is consistent with previous data indicating their involvement in the transduction of sour and high-salt stimuli. Sour is thought to be an indication of the freshness of a food source; rotting foods are rife with acid-producing microorganisms (DeSimone et al., 2001). Salt homeostasis is crucial for survival, so detecting and avoiding sources of damaging excess salt is likewise important. It follows that the activation of cells that respond to these qualities would induce avoidance behaviors. The degree to which our Pkd2l1-Cre mice avoided blue light in comparison with amber light was quite modest—a feature that may be due to experimental limitations or might indicate the particular role of the anterior tongue in taste-related behavior. Our Pkd2l1-Cre driver drove ChR2 expression in most, but not all PKD2L1 type III cells, which are less populous in the anterior tongue (Yoshida et al., 2009). Since the circumvallate taste buds of the posterior tongue are more difficult to stimulate with light, it is unlikely that our stimulation paradigm successfully activated as many posterior type III cells as anteriorly situated ones. Alternatively, the modest avoidance we see may indicate the role of anterior tongue in behavioral responses to tastants. Several lines of evidence suggest that the anterior tongue may be important for discriminating and

Figure 5. Pkd2l1-Cre, ChR2 mice do not increase licking behavior in response to light in the absence of water. A, Lick patterns during the first minute of behavioral experiments with 10 littermate control (gray) and 10 Pkd2l1-Cre, ChR2 (black) mice featuring a water spout without water or light (left), without water but with blue light (center), or with water and without light (right). B, Left, Total licks in the first minute of the experiments in A. For Pkd2l1-Cre, ChR2 mice (black), total licks did not differ between the water– light– condition and the water– light+ condition (t = 1.135, p = 0.2712, unpaired t test). Lick totals in both of these conditions were significantly different from the lick count for the water+ light– condition (t = 13 and t = 14, respectively; ****p < 0.0001 for each, unpaired t test). Right, Cumulative lick total over 10 min for Pkd2l1-Cre, ChR2 mice under each condition. All error bars denote 95% confidence intervals.
identifying specific tastants, while the posterior tongue acts more as a binary gatekeeper (Travers et al., 1987; Frank, 1991; Spector and Grill 1992; Hellekant et al., 1997; St John and Spector 1998; Tomchik et al., 2007; Yoshida et al., 2009). Anterior tongue, therefore, may not communicate aversion so much as stimulus identity.

Our behavioral data differ considerably from recent findings, which suggest that PKD2L1⁺ cells are water detectors as well as sour detectors (Zocchi et al., 2017). Recent studies observe that type III taste cells are more heterogeneous than previously assumed (Wilson et al., 2017; Lossow et al., 2017). This underlying molecular diversity within the type III cell population is unlikely to cause such divergent results as seen between this work and that of Zocchi et al. (2017), because both studies targeted the same genetic subpopulation. However, while both studies use Pkd2l1 to drive ChR2 expression, the tactics are slightly different; Zocchi et al. (2017) used a BAC transgenic mouse, which drives Cre expression from the Pkd2l1 promoter region, while we used a mouse featuring an IRES-Cre knock-in construct directly following the Pkd2l1 gene. These separate approaches may give rise to differences in expression levels and/or patterns, both of which might affect behavioral responses to light stimulation of the oral cavity. Our mouse, for example, drives ChR2 expression in only about half of the PKD2L1-immunoreactive cells. Perhaps a subtype of PKD2L1⁺ type III cells that was not activated in our experiments is responsible for water detection, and the behaviors observed by Zocchi et al. (2017) are the result of activation of this distinct population. ChR2 expression level differences between the two mouse models might also affect the level of activation of type III cells. Increased amounts of ChR2 in a single cell might increase the baseline electrical excitability on account of some constitutive activity of the channels. Differential type III cell excitability may, in turn, alter signaling to afferent nerve fibers. The synapse between type III cells and afferent nerve fibers has thus far been treated as a sort of “on/off” switch, but the nervous system often uses more nuanced methods of information transfer. Neurons involved in lower-frequency sound localization in gerbils, for example, code information not by maximal firing rate, but by a graded slope of firing rates (for review, see Ashida and Carr, 2011), and retinal bipolar cells generally function as integrators of currents without firing action potentials (for review, see Nelson and Connaughton, 2007). Perhaps coding at the type III cell–nerve synapse depends in part on the degree of depolarization, allowing for a more complicated transmission paradigm that may differ between our mouse models. Ultimately, more experimentation is necessary to tease out the causes for our differing results.

The data presented here demonstrate that direct activation of PKD2L1⁺ type III cells by light is sufficient to elicit responses in nerves innervating the anterior and posterior tongue, as well as aversive behavioral responses. Continued experimentation with this mouse model is necessary to determine the perceived taste quality of this optogenetic stimulation and to resolve the apparent conflicts with published data.

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