

Bitter triggers acetylcholine release from polymodal urethral chemosensory cells and bladder reflexes

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Chemosensory cells in the mucosal surface of the respiratory tract (“brush cells”) use the canonical taste transduction cascade to detect potentially hazardous content and trigger local protective and aversive respiratory reflexes on stimulation. So far, the urogenital tract has been considered to lack this cell type. Here we report the presence of a previously unidentified cholinergic, polymodal chemosensory cell in the mammalian urethra, the potential portal of entry for bacteria and harmful substances into the urogenital system, but not in further centrally located parts of the urinary tract, such as the bladder, ureter, and renal pelvis. Urethral brush cells express bitter and umami taste receptors and downstream components of the taste transduction cascade; respond to stimulation with bitter (denatonium), umami (monosodium glutamate), and uropathogenic *Escherichia coli*; and release acetylcholine to communicate with other cells. They are approached by sensory nerve fibers expressing nicotinic acetylcholine receptors, and intraurethral application of denatonium reflexively increases activity of the bladder detrusor muscle in anesthetized rats. We propose a concept of urinary bladder control involving a previously unidentified cholinergic chemosensory cell monitoring the chemical composition of the urethral luminal microenvironment for potential hazardous content.

Mucosal surfaces of the mammalian respiratory and gastrointestinal tract contain solitary epithelial cells with characteristic microvilli at their tip, from which the name “brush cells” derives (1–3). In the respiratory tract, these brush cells serve as sentinels, using the canonical taste transduction cascade to monitor the mucosal lining fluid for potential harmful substances, such as “bitter” bacterial products, and evoking reflexes aimed at combating further ingress of such compounds, such as closure of ducts leading into adjacent compartments (vomeronasal organ) or respiratory reflexes (4–7).

In line with such a sentinel function, brush cell abundance decreases with increasing distance to the opening to the outside world, being numerous in the nose and nearly absent in the intrapulmonary airways. Physiologically, the urinary tract allows passage in only one direction to release urine, but ascending infection by uropathogenic bacteria is not uncommon and is a major risk factor in fatal kidney disease and male infertility (8, 9). We hypothesized that the urogenital tract also may be equipped with chemosensory sentinel cells that monitor the lumen for potential hazardous content.

Results

Brush Cells Are Positioned at the Portal of Entry into the Urinary Tract. The majority of chemosensory brush cells of the mouse airways use acetylcholine as a signaling molecule and can be readily identified in mice expressing GFP driven by the promoter of the acetylcholine-synthesizing enzyme choline acetyltransferase (ChAT-eGFP mice) (4, 6). On screening of tissue sections

and whole-mount preparations of the urogenital tract of two independently generated ChAT-eGFP mouse strains (10, 11), we observed solitary ChAT-eGFP⁺ cells in the epithelium of the urethra, but not in the epithelial linings of the renal pelvis, ureter, urinary bladder, and pelvic segment of the vas deferens (Fig. 1 and Fig. S1). The total numbers of such urethral cells did not differ significantly between male and female animals (mean ± SD, 379 ± 99 and 551 ± 253, respectively, $P = 0.26$, t test; $n = 6$ for each sex). In male prostate, coagulating, and seminal glands and in both male and female paraurethral glands, these cells are located in the excretory ducts close to the opening into the urethra, but not in the glandular bodies themselves (Fig. S2). These cells vary in shape from the typical flask-like structure of tracheal brush cells, with a broad base at the basement membrane and an elongated tip reaching the lumen, to more complex morphologies with slender foot processes reaching the basal lamina directly or in an oblique course (Figs. 2 and 3). Horizontally oriented cell bodies with unclear connections to the luminal surface are seen as well (Fig. 3 *A* and *B*).

Double-labeling procedures revealed that these cells are distinct from the previously known urethral neuroendocrine cell population (12, 13) expressing protein gene product 9.5, serotonin, and chromogranin A (Fig. 2 *B* and *C* and Table S1). In

Significance

We report the presence of a previously unidentified cholinergic, polymodal chemosensory cell in the mammalian urethra, the potential portal of entry for bacteria and harmful substances into the urogenital system. These cells exhibit structural markers of respiratory chemosensory cells (“brush cells”). They use the classical taste transduction cascade to detect potential hazardous compounds (bitter, umami, uropathogenic bacteria) and release acetylcholine in response. They lie next to sensory nerve fibers that carry acetylcholine receptors, and placing a bitter compound in the urethra enhances activity of the bladder detrusor muscle. Thus, monitoring of urethral content is linked to bladder control via a previously unrecognized cell type.

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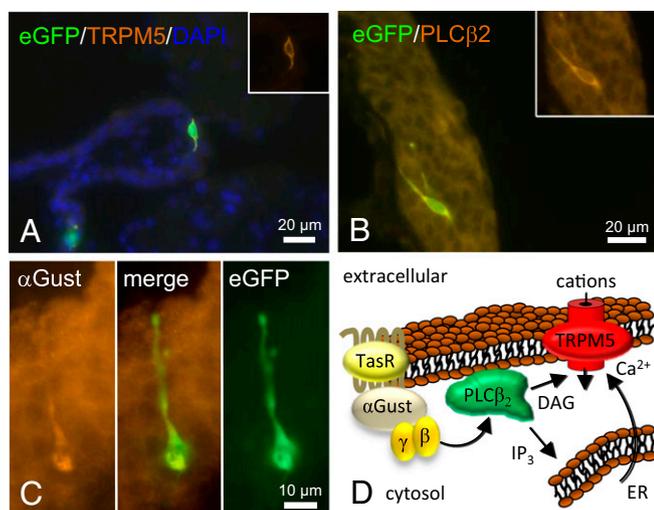


Fig. 3. Cholinergic urethral brush cells express components of the canonical taste transduction cascade. (A) Male urethra. ChAT-eGFP (green) and TRPM5 (orange) are expressed by the same cell; nuclei are labeled in blue with DAPI. (B) Female urethra. An obliquely oriented ChAT-eGFP cell is immunolabeled for PLC β 2. In this section plane, no connection of this cell to the urethral lumen is visible. (C) Male urethra, diverticular region. α -Gustducin (α Gust)⁺ ChAT-eGFP cell extending a slender process toward the lumen. (D) Schematic drawing of the canonical transduction cascade known from taste buds (based upon information published in ref. 49) depicting the position and role of α -gustducin, PLC β 2, and TRPM5 in this process. DAG, diacylglycerol; ER, endoplasmic reticulum; IP $_3$, inositol trisphosphate; TasR, taste receptor.

In addition, immediate and highly reversible currents were triggered by short, repetitive pulses of denatonium (Fig. S4).

Cholinergic Urethral Brush Cells Are Polymodal Chemosensors. Tas1R family members (Tas1R1–3) participate in sweet and umami (free L-amino acids) perception. Isolated urethral cholinergic chemosensory cells expressed mRNAs coding for Tas1R1 and Tas1R3, whereas Tas1R2 expression was not detected (Fig. 4A). Tas1R1–Tas1R3 coexpression yields an umami receptor (20). Accordingly, urethral brush cells responded with increases in [Ca²⁺]_i to monosodium glutamate (umami, 25 mM; 90%; 38/42 cells) in addition to denatonium and ATP (0.5 mM; 91%; 43/47 cells). The majority of cells (86%; 36/42) responded to both the bitter receptor agonist denatonium and glutamate, exhibiting properties of a polymodal (bitter/umami) chemosensor (Fig. 4F).

Notably, not all bitter substances triggered responses in urethral brush cells. In line with the lack of detectable Tas2R105 mRNA in urethral brush cells (Fig. 4A), cycloheximide, a Tas2R105 agonist, did not evoke changes in [Ca²⁺]_i at a concentration of 0.1 mM (Fig. 4G), which is known to stimulate a [Ca²⁺]_i increase in Tas2R105-transfected cells (21). In addition to synthetic bitter and umami stimuli, heat-inactivated uropathogenic *Escherichia coli* (strain CFT073; 2–5 × 10⁷ cfu) also triggered a rise in [Ca²⁺]_i (Fig. 4H). Tas1R2–Tas1R3 coexpression yields a sweet receptor (22). Consistent with the absence of detectable Tas1R2 mRNA in urethral brush cells (Fig. 4A), the artificial sweetener saccharin (5 mM) did not trigger a [Ca²⁺]_i response in urethral brush cells (Fig. 4G).

Urethral Brush Cells Use Acetylcholine for Stimulus-Induced Paracrine Signaling. The concept of brush cells serving as chemosensory sentinels monitoring the chemical composition of mucosal surface fluids and initiating local or reflexive responses implies stimulus-induced signaling to neighboring cells. In view of these cells' ChAT expression, we tested for acetylcholine as a potential messenger released by urethral brush cells. Acetylcholine content in the supernatant (0.5 mL) of isolated urethral cells was increased by an average of 0.875 pmol (from a baseline of 10.0 pmol)

in response to stimulation with denatonium (25 mM, 5 min) (Fig. 5A). In our experiments, we noted a rise in [Ca²⁺]_i in response to denatonium in non-GFP-expressing cells when situated in the vicinity of ChAT-eGFP⁺ cells, but not in isolated cells resting on coverslips devoid of ChAT-eGFP cells (Fig. S5). Such cells express functional acetylcholine receptors; administration of 25 μ M acetylcholine together with the esterase inhibitor physostigmine (5 μ M) evoked a rise in [Ca²⁺]_i in eGFP⁻ cells that was sensitive to a cholinergic blocker mixture (2 μ M atropine and 20 μ M mecamylamine) (Fig. 5B). In the presence of the same cholinergic blockers, denatonium still evoked a rise in [Ca²⁺]_i, even at a slightly enhanced level, in ChAT-eGFP cells, but no longer in eGFP⁻ cells, demonstrating stimulus-evoked cholinergic signaling between chemosensory cells and surrounding cells (Fig. 5C).

Sensory Nerve Fibers Approach Urethral Brush Cells and Elicit Reflex Bladder Activation on Urethral Bitter Substance Application.

Cholinergic chemosensory cells of the respiratory tract are approached by cholinergic sensory nerve fibers that initiate protective respiratory reflexes (4–6). The dominant nicotinic acetylcholine receptor (nAChR) subtype of viscerosensory neurons contains the α 3 subunit, and such neurons project to the murine lower urinary tract (23). Using a GFP reporter mouse strain for this promoter (24), we identified a dense nerve fiber network immediately underneath and partially penetrating into the urethral epithelium, coming into contact with villin-positive brush cells (Fig. 6A). Because urethral sensory nerve fibers are known to be linked to the micturition reflex and to coordinate muscle constriction of the bladder and urethra (25), we assessed for reflex coupling between urethral bitter sensing and detrusor activity. The mouse urethra is too small to allow simultaneous cystometric recording of intravesical pressure and urethral manipulation, and thus rat is the model of choice for this purpose (26).

We first validated by immunohistochemistry the occurrence of solitary villin-, α -gustducin-, PLC β 2-, TRPM5-, and ChAT-immunoreactive cells in the urethral mucosa of rats (Fig. 6B). Notably, such cells were also detected in two human urethral specimens obtained during prostatectomy (Fig. 6B), demonstrating that their occurrence is not restricted to rodents. Continuous filling of the bladder with saline solution (0.04 mL/min) through a catheter inserted into the bladder dome simulated natural bladder filling in fast motion and caused cycles of rising intravesical pressure and micturition in urethane-anesthetized rats. Urethral instillation of saline solution (0.9%; 50 μ L) through the external urethral orifice augmented pressure rises only slightly, but a single dose of denatonium (25 mM; 50 μ L) significantly increased detrusor activity far beyond the first micturition, causing washout or a drastic dilution of urethral content (Fig. 6C and D). Lower denatonium concentrations (2.5–12.5 mM; $n = 2$ each) had no obvious effect, resembling the effectiveness of denatonium on [Ca²⁺]_i rises in isolated urethral chemosensory cells. Intraurethral administration of the general nicotinic receptor blocker mecamylamine (10⁻⁴ M) significantly reduced the denatonium-induced increase in detrusor activity, although the activity did not completely return to baseline (Fig. 6C and E). When administered before denatonium, mecamylamine also did not completely block the denatonium-induced rise in detrusor activity ($n = 2$).

Discussion

Up to now, solitary chemosensory or brush cells have been identified in the respiratory and gastrointestinal tracts, but not in any other mammalian organ system. Likely owing due to their anatomic restriction to the portal of entry into the urogenital tract (i.e., the urethra and glandular ducts opening into it), these chemosensory cells have escaped detection in previous searches for urogenital brush cells that focused on more centrally located organs, including the kidney, uterus, and prostate gland, before the sentinel function of solitary chemosensory cells had been proposed (14). Coexpression patterns of various components of

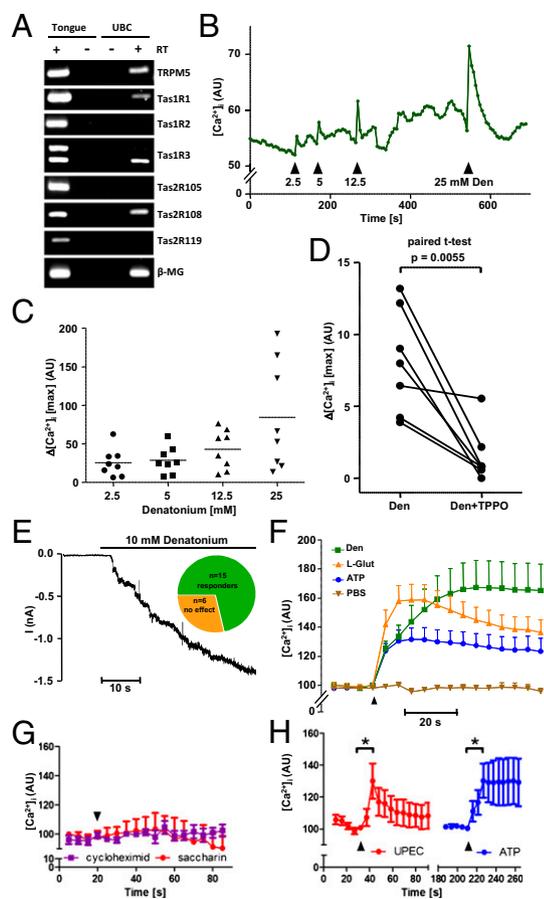


Fig. 4. Urethral cholinergic brush cells are polymodal chemosensors. (A) RT-PCR, agarose gel. Urethral brush cells (UBCs) were isolated by magnetic beads coated with TRPM5 antibody. Tongue, positive control; β 2-microglobulin (β -MG), housekeeping gene; +/- RT, aliquots processed with/without reverse transcription. (B and C) Dose-dependent increase in $[Ca^{2+}]_i$ in dissociated UBC in response to the Tas2R108 agonist denatonium. All drugs were added under continuous flow in the chamber, so that indicated concentrations were reached initially and then washed out. (B) Confocal laser scanning recording of changes in Calcium Orange fluorescence in a single cholinergic (eGFP⁺) brush cell. The y axis depicts arbitrary units (AU) correlating to $[Ca^{2+}]_i$. (C) The y axis depicts maximum $[Ca^{2+}]_i$ increases in eight cholinergic cells in response to increasing denatonium concentrations. (D) Same technique as in B and C. Shown are maximum $[Ca^{2+}]_i$ increases evoked by 25 mM denatonium before and after application of the TRPM5 inhibitor TPPO (0.25 mM) in seven cholinergic brush cells. (E) Whole-cell patch-clamp recording, with membrane potential clamped to -60 mV. Increased inward current was observed in 15 of 21 cells (responders). (F–H) Same technique as in B–D. (F) Superimposed calcium increases in 32 polymodal cells that responded to all three stimuli (0.5 mM ATP, 25 mM denatonium, and 25 mM L-glutamate). (G) In contrast, dissociated cells did not respond to the Tas2R105 agonist cycloheximide (Cyc; 0.1 mM) or to the artificial sweetener saccharin (Sac; 5 mM). (H) Heat-inactivated uropathogenic *E. coli* (UPEC; 2.5×10^7 cfu) induced $[Ca^{2+}]_i$ increase to the same extent as ATP. Graphs depict mean and SEM. * $P < 0.05$, paired *t* test.

the taste transduction cascade and ChAT-eGFP suggest the presence of more than one chemosensory cell type in the urethra, including a cholinergic type using PLC β 2 and TRPM5, proteins essential for oropharyngeal bitter and umami perception (27), for downstream signaling. Consistent with our $[Ca^{2+}]_i$ measurements, this signaling pathway involves G $\beta\gamma$ stimulation of PLC β 2 with subsequent release of Ca^{2+} from intracellular stores (28).

The role of the taste-specific G α protein α -gustducin (29), which is expected to activate a phosphodiesterase, is less clear (28). In taste buds, α -gustducin is inconsistently coexpressed with

T1R1–3, particularly in the posterior tongue (30), and its genetic ablation diminishes, but does not abrogate, bitter and umami gustation (31, 32). Similarly, the vast majority ($\geq 90\%$) of cholinergic urethral brush cells respond to bitter (denatonium) and umami (L-glutamate), but only approximately one-third coexpress α -gustducin.

A striking feature of cholinergic/TRPM5⁺ urethral chemosensory cells is the coexpression of taste receptors of the Tas1R and Tas2R families and the resulting polymodal response to both bitter and umami stimuli. Similarly, solitary nasal chemosensory cells coexpress Tas1R3 with Tas2R5 and Tas2R8 (33); however, although the responses of these cells to bitter stimuli are well documented (34), whether they also respond to umami remains to be established. This situation is in contrast to oropharyngeal gustation, where Tas1R and Tas2R receptors are not coexpressed in taste buds (33, 35), and 83% of receptor cells in murine vallate papillae respond to only a single taste quality, which provides a basis for taste coding in taste buds (36).

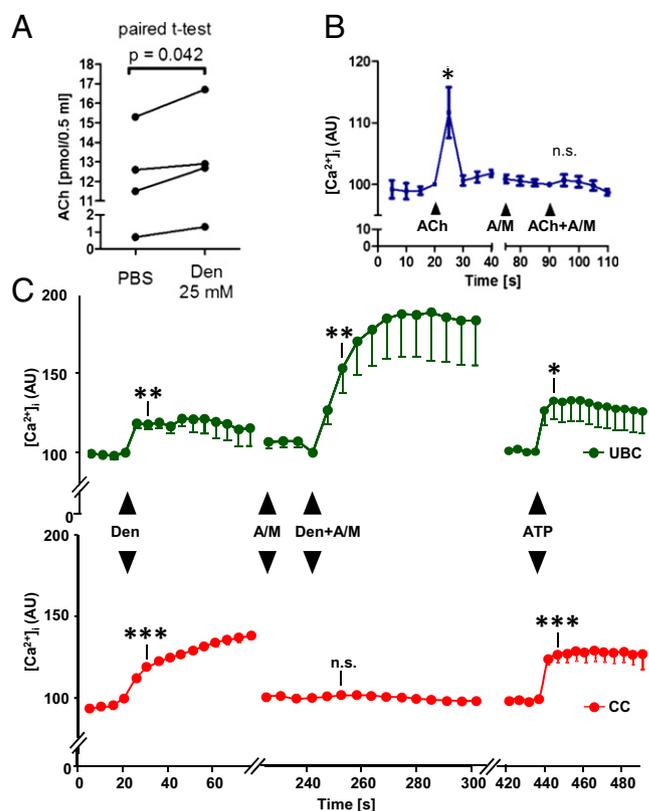


Fig. 5. Urethral brush cells release acetylcholine. (A) Acetylcholine (ACh) content in the supernatant (0.5 mL) of isolated urethral cells after a 5-min exposure to PBS (control) or denatonium. Isolates from one urethra are treated as paired data. (B) Calcium recordings from eight eGFP⁺ urethral cells located on coverslips that did not contain eGFP⁺ cells. In the continuous presence of the acetylcholinesterase inhibitor physostigmine (5 μ M), acetylcholine (25 μ M) evokes an increase in $[Ca^{2+}]_i$ that is sensitive to a mixture (A+M) of muscarinic (2 μ M atropine) and nicotinic (20 μ M mecamylamine) acetylcholine receptor blockers. (C) Parallel $[Ca^{2+}]_i$ recordings from eGFP⁺ ($n = 13$, Upper) and eGFP⁻ ($n = 74$; Lower) isolated urethral cells located in their vicinity (same field of view during confocal laser scanning recording) on the coverslip. Both cell types respond to denatonium (Den; 25 mM) with a $[Ca^{2+}]_i$ increase. This responsiveness is lost in eGFP⁻ cells, but persists (even at enhanced levels; $P = 0.04$, paired *t* test) in eGFP⁺ cells after pretreatment with cholinergic blockers (A+M; A), demonstrating denatonium-evoked cholinergic signaling from eGFP⁺ to eGFP⁻ cells. Response to the nongustatory stimulus ATP demonstrates the viability of cells at the end of the experiment. n.s., nonsignificant. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$, paired *t* test compared with value immediately before substance application.

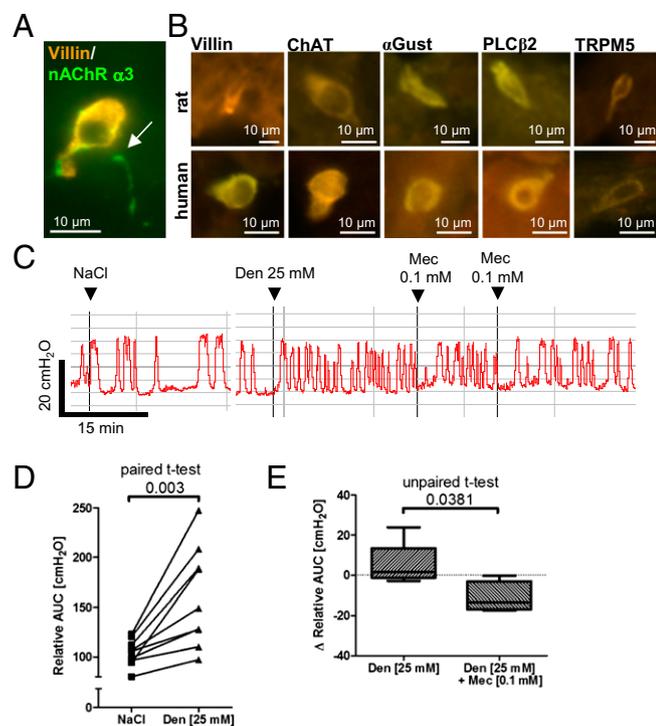


Fig. 6. Sensory nerve fibers approach urethral brush cells and elicit reflex bladder activation on application of a bitter substance. (A) A sensory nerve fiber expressing eGFP under the control of the nicotinic receptor $\alpha 3$ -subunit (arrow) establishes contact with a villin-immunoreactive brush cell in the mouse urethra. (B) Immunolabeling. Solitary epithelial cells with immunoreactivity to brush cell markers, taste transduction cascade components, and ChAT in rat and human urethral epithelium. (C–E) Cystometric recordings from urethane-anesthetized rats. The bladder was continuously filled with saline (0.04 mL/min), causing a rise in intravesical pressure and initiating detrusor contraction and micturition. (C) Original recording of a single experiment showing that urethral instillation of denatonium increases detrusor activity compared with saline application, which is diminished by the nicotinic blocker mecamylamine (Mec). (D and E) Detrusor activity quantified as AUC. (D) Denatonium increases detrusor activity. AUC was determined over a period of 18–36 min in each condition, in nine experiments. (E) First recordings were made after stimulation with a single dose of denatonium for a minimum of 18 min, with AUC/min as the baseline value. AUC/min decreased after application of mecamylamine, but increased over time when no additional compound was applied.

In oropharyngeal gustation, bitter represents an aversive stimulus and umami represents a rewarding stimulus, raising the question as to the possible functional meaning of sensing of both qualities by a single cell. In contrast, on other mucosal surfaces, such as the urethral lining, these qualities represent potentially harmful (aversive) content. Bacteria produce and secrete bitter receptor-activating substances (5, 37, 38). In biofilms, such substances from the Gram-negative bacterium *Pseudomonas aeruginosa*, one of the predominant causative microorganisms in catheter-associated urinary tract infection (39), can reach concentrations as high as 600 μM (40). On the other hand, glutamate metabolism is positively linked to the pathogenic potential of *Proteus mirabilis* in the urinary tract (41), and free amino acids (i.e., umami) facilitate bacterial growth in urine (42). Thus, the bitter/umami polymodality of chemosensory cells may serve to broaden the spectrum for recognition of potential hazardous material in the urethral lumen. Most importantly, these chemosensory cells responded to heat-inactivated uropathogenic *E. coli*, the primary cause of urinary tract infection (43).

Acetylcholine, a secretory product of these cells, may alter sensitivity of this process in an autocrine manner, as demonstrated by the enhanced tastant response after application of a

muscarinic/nicotinic receptor blocker mixture. In the more complex lingual taste buds, autocrine cholinergic signaling enhances taste signaling via muscarinic receptors (44). Our cell culture experiments revealed that the amount of acetylcholine released on bitter stimulation is also sufficient to induce paracrine effects. In situ, these cholinergic cells, like those in the trachea (6), are directly approached by sensory nerve fibers expressing the nicotinic acetylcholine receptor $\alpha 3$ subunit, and luminal denatonium evoked reflex activation of the detrusor muscle at a concentration that stimulated urethral chemosensory cells. This reflex activation was sensitive to local application of a nicotinic receptor blocker, demonstrating the involvement of cholinergic, nicotinic transmission from chemosensory cells to sensory nerve fibers. Nonetheless, the denatonium-induced increase in detrusor activity was not entirely abrogated, possibly owing to (i) additional involvement of excitatory muscarinic acetylcholine receptors that are also expressed by urinary tract afferent neurons (45); (ii) additional involvement of a cotransmitter, such as ATP, which transmits information from taste cells to afferent fibers in taste buds (46); or (iii) insufficient access of intraluminally applied mecamylamine to the basolaterally located communication site between chemosensory cells and nerve fibers, as lower urinary tract epithelia form an extraordinary tight barrier (47).

In conclusion, we propose a concept of urinary bladder control involving a previously unidentified cholinergic chemosensory cell monitoring the chemical composition of the urethral luminal microenvironment for potential hazardous content.

Materials and Methods

Animals. Two independently generated ChAT^(BAC)-eGFP mice were provided by M. Kotlikoff (Cornell University), and H. Monyer (University of Heidelberg) (10, 11). Tg(*Chrna3*-EGFP)BZ135Gsat mice with eGFP driven by the nAChR $\alpha 3\beta 4\alpha 5$ cluster were provided by I. Ibanez-Tallon (MDC Molecular Medicine) (24). C57BL/6 mice were obtained from The Jackson Laboratory. Male Crl:WI Wistar rats were obtained from Charles River Deutschland. All animals were housed under standard laboratory conditions (12 h dark, 12 h light). Mice were killed by inhalation of an overdose of isoflurane (Abbott) and exsanguination. The experiments were approved by the local authorities (Rp Giessen, Germany; reference nos. A9/2011, A11/2011, A60/2012, A61/2012, and 12/2013).

Cell Isolation. Urethrae were dissected, cut into small pieces, and enzymatically digested in dispase (2 mg/mL; Sigma-Aldrich) for 30 min in HBSS (Invitrogen) and 5 min in trypsin/PBS (1:1; Invitrogen) at 37 °C. After centrifugation (60 \times g, 5 min) and mechanical dissociation, cells were resuspended in PBS and separated through a cell strainer (70 μm ; BD Bioscience). Chemosensory cells were identified by either eGFP or a rabbit polyclonal TRPM5-antibody (ab72151, 1:125; Abcam) directed against an extracellular domain. Cells were incubated for 1 h at 37 °C with this primary antibody, followed by a 1-h incubation with FITC-conjugated donkey anti-rabbit IgG (1:125; Millipore) at 37 °C. The same procedure, but using Cy5-conjugated donkey anti-rabbit IgG (1:400; Dianova), was also applied to cells isolated from ChAT-eGFP mice. As observed in the immunohistochemistry of tissue sections (Table S3), TRPM5 labeling and eGFP expression matched nearly 1:1. For subsequent RT-PCR analysis (primer sequences given in Table S4), chemosensory cells were isolated by incubating dissociated urethra first with the TRPM5 antibody as described above and then with magnetic beads (Invitrogen) coated with goat anti-rabbit IgG (H+L) (PI65-6100; Invitrogen), followed by harvesting by magnetic cell separation.

Measurement of Intracellular Calcium Concentration. Isolated cells were loaded with fluorescent calcium indicator Calcium Orange AM (3 μL) in Tyrode III solution (297 μL ; 8 mM CaCl_2 , 130 mM NaCl, 5 mM KCl, 1 mM MgCl_2 , 10 mM HEPES, 10 mM glucose, 10 mM pyruvic acid, and 5 mM NaHCO_3) according to the manufacturer's protocol (Invitrogen) and then plated on coverslips for 30–60 min at 37 °C. Intracellular calcium concentration was analyzed with a confocal laser scanning microscope (Zeiss LSM 710; 561-nm wavelength generated by a DPSS 561-10 laser) during continuous superfusion (3 mL/min) with Tyrode solution. Fluorescence intensities at the start of the recording period were set arbitrarily at 100%. Test stimuli and concentrations were adenosine 5'-triphosphate bis(Tris) salt dihydrate (ATP, 0.5 mM; Sigma-Aldrich), cycloheximide (0.1 mM; Sigma-Aldrich), denatonium benzoate (25 mM; Molekula), L-glutamic acid monosodium salt monohydrate (L-glutamate, 25 mM; Sigma-Aldrich),

saccharin (5 mM; Fluka), acetylcholine chloride (25 μ M; Sigma-Aldrich), and heat-inactivated uropathogenic *E. coli* [UPEC strain CFT073 (NCBI: AE014075, NC_004431), $\sim 2\text{--}5 \times 10^7$ cfu, provided by T. Chakraborty, JLU Giessen]. Inhibitors were TPPO (0.25 mM, Sigma-Aldrich), eserine hemisulfate (10 μ M; Sigma-Aldrich), mecamlamine hydrochloride (0.02 mM; Sigma-Aldrich), and atropine sulfate (0.002 mM; RBI). Data are presented as mean \pm SEM and were analyzed by the two-tailed paired *t* test.

Patch-Clamp Recordings. Cells were isolated from ChAT-eGFP mice as described for $[Ca^{2+}]_i$ recordings, and eGFP-expressing cells were identified with a fluorescence microscope (Zeiss Axiovert 10). The bath solution consisted of 140 mM NaCl, 4.5 mM KCl, 2.5 mM $CaCl_2$, 1 mM $MgCl_2$, 10 mM Hepes, and 5 mM D -glucose (pH 7.4). The pipette solution consisted of 10 mM NaCl, 18 mM KCl, 92 mM K-gluconate, 0.5 mM $MgCl_2$, 1 mM EDTA, and 10 mM Hepes (pH 7.2). The liquid junction potential was absorbed with the clamped voltage, producing an effective membrane potential of -60 mV. The bath solution included appropriate amounts of mannitol to compensate for osmotic changes or different concentrations of denatonium (1–25 mM), applied by a pressure-driven perfusion system. Transmembrane currents were amplified (EPC 9; Heka Electronics) and recorded continuously (sampled with 10 kHz, filtered with 3 kHz) with Pulse 8.77 software (Heka Electronics).

Urodynamic Measurement. Rats were anesthetized by an s.c. injection of urethane (1.2 g/kg) at 1 h before surgery and maintained under anesthesia during the surgical preparation while on a heating pad (37 $^{\circ}C$). A catheter (PE 50; Intramedic) was inserted into the bladder dome and connected to a pressure transducer and an infusion pump. Saline solution at room temperature was infused into the bladder at a rate of 0.04 mL/min. After a stabilization phase of 15–30 min, the intravesical bladder pressure was recorded continuously, and 50 μ L of test stimuli were delivered into the urethral external orifice via a 0.9×25 mm cannula (Braun Vasofix G22) mounted on a 100- μ L pipette. For final data analysis, areas under the curve (AUC) of equal time periods before and after stimulation were compared; data are presented as AUC/min. The urodynamic recording sessions took 3–4 h for each animal.

Fluorescence microscopy and immunohistochemistry, pre-embedding immunohistochemistry, and EM, RT-PCR, and acetylcholine measurements were performed as described previously (6, 48), with minor modifications. More details are provided in *SI Materials and Methods*.

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