Intraluminal acid activates esophageal nodose C fibers after mast cell activation

Shizhong Zhang, Zhenyu Liu, Andrea Heldsinger, Chung Owyang, and Shaoyong Yu

1Division of Gastroenterology and Hepatology, Department of Medicine, University of Michigan Medical School, Ann Arbor, Michigan; and 2Johns Hopkins University School of Medicine, Baltimore, Maryland

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Zhang S, Liu Z, Heldsinger A, Owyang C, Yu S. Intraluminal acid activates esophageal nodose C fibers after mast cell activation. Am J Physiol Gastrointest Liver Physiol 306: G200–G207, 2014. First published November 21, 2013; doi:10.1152/ajpgi.00142.2013.—Acid reflux in the esophagus can induce esophageal painful sensations such as heartburn and noncardiac chest pain. The mechanisms underlying acid-induced esophageal nociception are not clearly understood. In our previous studies, we characterized esophageal vagal nociceptive afferents and defined their responses to noxious mechanical and chemical stimulation. In the present study, we aim to determine their responses to intraluminal acid infusion. Extracellular single-unit recordings were performed in nodose ganglion neurons with intact nerve endings in the esophagus using ex vivo esophageal-vagal preparations. Action potentials evoked by esophageal intraluminal acid perfusion were compared in naive and ovalbumin (OVA)-challenged animals, followed by measurements of transepithelial electrical resistance (TEER) and the expression of tight junction proteins (zona occludens-1 and occludin). In naive guinea pigs, intraluminal infusion with either acid (pH 2–3) or capsaicin did not evoke an action potential discharge in esophageal nodose C fibers. In OVA-sensitized animals, following esophageal mast cell activation by in vivo OVA inhalation, intraluminal acid infusion for about 20 min started to evoke action potential discharges. This effect is further confirmed by selective mast cell activation using in vitro tissue OVA challenge in esophageal-vagal preparations. OVA inhalation leads to decreased TEER and zona occludens-1 expression, suggesting an impaired esophageal epithelial barrier function after mast cell activation. These data for the first time provide direct evidence of intraluminal acid-induced activation of esophageal nociceptive C fibers and suggest that mast cell activation may make esophageal epithelium more permeable to acid, which subsequently may increase esophageal vagal nociceptive C fiber activation.

Acid reflux in the esophagus can induce esophageal painful sensations such as heartburn and noncardiac chest pain. Proton pump inhibitor (PPI) therapy is usually effective for the symptoms after PPI therapy (23). A better understanding of the mechanisms of acid-evoked esophageal nociception will be helpful to develop an effective pain modulator to manage these symptoms in addition to PPIs. Currently, the underlying mechanism of acid-evoked activation of esophageal nociceptor is largely unknown (1). Recent studies have shown that several ion channels or receptors in sensory neurons are responsible for sensing acid (11). These studies provide important information on acid-evoked responses from sensory neurons, the cell bodies of sensory nerves, but the knowledge on acid-evoked responses of sensory nerve endings in peripheral tissues is still lacking.

Sensory nerves detect stimuli resulting from the physiological activity of the tissue as well as stimuli associated with impending and/or actual tissue damage (noxious stimuli). The sensory nerves capable of discriminating noxious stimuli are termed nociceptors (32). The transduction of noxious stimulation in the esophagus starts from sensory nerve terminals in the wall of the esophagus. The evoked action potential discharge is conducted through both vagal and spinal pathways into the central nervous system to generate nociception (3). Our previous studies demonstrated that esophageal vagal afferents have distinctive subtypes of nociceptive C fibers, namely nodose and jugular C fibers. They are able to discriminate noxious and innocuous esophageal distensions and respond to noxious chemical (capsaicin) stimulation (32). Whether these nociceptive C fibers are able to sense intraluminal acid has yet to be determined.

The normal esophageal epithelial barrier is usually resistant to acid exposure (6, 8). But acid combined with pepsin or bile salts can disrupt the esophageal epithelial barrier (6, 27, 28). Impaired mucosal barrier function may allow intraluminal acid to reach the sensory nerve endings underneath, activating those nociceptive afferents and inducing esophageal nociception (1). It has been reported that mast cell activation damages the intestinal epithelial barrier (18, 22, 24, 26). But the effect of mast cell activation on esophageal epithelial barrier function has not been reported. Our previous studies have established models of esophageal mast cell activation induced by either in vitro tissue antigen using ovalbumin (OVA) challenge (29) or in vivo antigen inhalation (31) in antigen-sensitized animals. Based on these present-day established models, we test the hypothesis that intraluminal acid activates esophageal vagal nodose C fibers after mast cell activation, due mainly to impaired esophageal epithelial barrier function.

METHODS

Male Hartley guinea pigs (Hilltop Laboratory Animals, Scottsdale, PA) weighing 100–300 g were used. All experiments were approved by the University of Michigan Committee on Use and Care of Animals (UCUCA, #PRO0000117).

Animal sensitization and challenge. According to our previous studies (29, 31), guinea pigs were sensitized with three intraperitoneal injections of OVA (10 mg/kg in 0.9% saline) every 48 h. Three weeks after the last injection, guinea pigs were ready for either in vitro tissue antigen challenge or in vivo antigen inhalation. For tissue antigen challenge, the dissected esophageal-vagal preparations from OVA-sensitized animals were immersed in Krebs bicarbonate solution (KBS) containing antigen (OVA, 10 μg/ml) for 30 min to induce antigen-mediated mast cell degranulation. For antigen inhalation, OVA-sensitized animals were exposed to aerosolized antigen (0.1%
OVA) in a plastic chamber for 1–10 min, depending on the development of dyspnea. The OVA was dissolved in 0.9% saline and delivered using a nebulizer driven by compressed air. The guinea pigs were closely monitored for signs of any allergic response such as gasping and increased respiratory rate. Once the guinea pigs developed such a response, they were removed and allowed to breathe ambient air. OVA inhalation was performed three times (once every 24 h) in each animal. Extracellular recordings were performed in ex vivo esophageal-vagal preparations after the last inhalation.

**Extracellular single-unit recording.** In previously described esophageal-vagal preparations, extracellular single-unit recordings of action potential discharges were performed from nodose ganglia neurons with intact C fiber nerve endings in the esophagus (32). Briefly, the esophagus and trachea were dissected with intact bilateral extrinsic vagal innervations (including jugular and nodose ganglia). The tissue was pinned in a small Sylgard-lined Perspex chamber filled with KBS at 35°C, and gassed with 95% O2-5% CO2. The chamber had two compartments: the esophagus with attached trachea and the vagus were pinned in the tissue compartment, and the vagus nerves including the nodose and jugular ganglia were pinned in the recording compartment. Isobaric esophageal distension for 20 s with an intraluminal pressure of 10–60 mmHg was used to determine the distension pressure-nerve activity relationship of an esophageal afferent fiber. Extracellular single-unit recordings were performed using an alumino-silicate glass microelectrode which was placed into a nodose ganglion with an electrode holder connected directly to the headstage (A-M Systems, Everett, WA). The recorded signal was amplified (microelectrode AC amplifier 1800, A-M Systems) and filtered (low cutoff, 0.5 kHz; high cutoff, 1 kHz), and the resultant activity was displayed on an oscilloscope (TDS 340, Tektronix, Beaverton, OR) and chart recorder. The data were stored and analyzed using the software TheNerveOfIt (sampling frequency 33 kHz; PHOCIS, Baltimore, MD).

We identified esophageal vagal afferent fibers according to our previously described methods (32). Briefly, the recording electrode was micromanipulated into the nodose or jugular ganglion (left or right). A distension-sensitive unit was identified when esophageal distension (in most experiments with a rapid increase in intraluminal pressure to 10 mmHg for 5 s) evoked action potential discharge. The serosal surface of the esophagus was then searched with a punctate mechanical probe (Von Frey hair, 1 mN, filament diameter <0.5 mm) applied to the tissue. A mechanosensitive receptive field was located when the punctate stimulus evoked discharge of action potentials with waveforms identical to the action potentials evoked by distension. The receptive field was then stimulated electrically (pulse duration 1 ms, frequency 1 Hz) with a concentric electrode inserted into the esophagus with the tip positioned at the site of the mechanosensitive receptive field. The initial voltage (100 V) was gradually reduced to the lowest voltage (threshold voltage) at which each stimulation pulse was followed by a single action potential (30–90 V for most afferent nerve fibers recorded). The waveforms of the electrically evoked action potentials were identical to those induced by distension and the punctate mechanical stimulus. Conduction time was measured as the time between the stimulation pulse and the action potential (visualized by oscilloscope).

The nerve fiber was considered a C fiber if it conducted action potentials at <1 m/s. Conduction velocity was calculated by dividing the length of the approximated nerve pathway (from the recorded nodose neurons to the mechanosensitive receptive field in the esophagus) by conduction time. The peaks of action potential discharges of nodose C fibers in response to different treatments or stimuli were analyzed and compared. The compounds used in the experiment included OVA (as antigen for animal sensitization and mast cell activation), HCl (pH = 2–3, adjusted with KBS), capsaicin [transient receptor potential vanilloid-1 (TRPV1) agonist, which usually activates C fibers], ATP, prostaglandin E2 (PGE2), serotonin, histamine, and bradykinin (all from Sigma-Aldrich, MO). The compounds were diluted in KBS to final concentration on the day of use.

**Histology.** Esophageal segments were fixed in 10% formaldehyde and embedded in paraffin. Sections of 5 μm were mounted on Superfrost Plus glass slides (Fisher Scientific) and stained with hematoxylin and eosin (H-E). The slides were analyzed under light microscope. The inflammation grade of the esophagus was evaluated according to a previously reported method (12), including assessments of active inflammation (neutrophil infiltration in the epithelium), the length of vascular papillae, basal-zone hyperplasia, and the number of intraepithelial eosinophils. In a separate study, esophageal segments from both naive (n = 3) and OVA-inhalation (n = 4) groups were fixed in Carnoy’s solution and embedded in paraffin. Sections of 6 μm were mounted on Superfrost Plus glass slides (Fisher Scientific) and stained with toluidine blue. The slides were analyzed under a light microscope, and the numbers of mast cells in the esophageal cross-sections were counted and compared.

**Transmucosal electrical resistance (TER).** The TEER of esophageal epithelium was measured as previously described (5). Following extracellular recordings, the mucosal epithelium of the esophagus was dissected and cut into three pieces (3.5 × 3.5 mm each). Each segment was sandwiched between two Plexiglas inserts with a 3-mm-diameter central hole, introduced into Costar snapwells, and then placed in the incubator (37°C, 5% CO2) for 30 min to stabilize the pH. The TEER was measured in the micro-snapwell system by using a planar electrode (Endom SNAP electrode attached to an Evom-G WPI analyzer, World Precision Instruments, Sarasota, FL). The result was averaged from the measurements of three segments from each esophagus and expressed in ohms per square centimeter.

**Western blot.** Equal amount of lysates (20 μg) freshly obtained from esophageal mucosal layers of naive (n = 4) and OVA-inhalation plus acid-infused (n = 6) animals were separated on Ready Gel 12% Tris-HCl, transferred to nitrocellulose Hybond enhanced chemiluminescence (ECL) membranes, and blotted with primary antibodies (overnight), and then secondary antibodies (1 h), followed by detection using ECL reagents (Pierce, IL). The membranes were exposed to ECL buffer for 30 s or 5 min and then high chemiluminescence film in the dark. The resulting bands were scanned and analyzed. The primary antibodies used in Western blot analysis included rabbit anti-zona occludens-1 (anti-ZO-1) antibody (1:1,000, #61–7300, Invitrogen, CA) and mouse anti-occludin antibody (1:500, #33–1500, Invitrogen, CA). The secondary antibody included goat anti-rabbit IgG horseradish peroxidase (HRP) (1:2,000, #sc-2004, Santa Cruz, CA), and goat anti-mouse-HRP (1:4,000, #sc-2005, Santa Cruz, CA).

**Data analysis.** In extracellular studies, we only analyzed the results from capsaicin-responsive nodose C fibers, which were confirmed by the end of each recording to indicate that the nerve terminals were exposed to chemical perfusion. We recorded afferent nerve activities from one nodose C fiber per animal, so the number of recorded fibers (n) equals the number of animals (N) used in the study. The chemical-evoked nerve response was quantified as peak frequency of the action potential discharge within a 5-min period, and averaged six recording periods for a total of 30 min. The peak frequency (Hz) of action potential discharges were presented as means ± SE and compared by one-way ANOVA. In the histological study, the inflammation grades from each group with different treatments were counted and compared by one-way ANOVA. In the permeability study, the TEERs were present as means ± SE and compared by one-way ANOVA. The density of each band on the Western blot was analyzed by densitometry, and normalized for quantitative comparisons by student’s t-test. For all experiments, significance was defined as P < 0.05.

**RESULTS**

In extracellular recordings using ex vivo esophageal-vagal preparations, a total of 84 esophageal nodose C fibers from 84 animals (one from each animal) was recorded to determine...
their responses to intraluminal acid infusion, with or without OVA sensitization plus OVA tissue challenge in vitro or OVA inhalation in vivo. These C fibers (with conduction velocity <1 m/s) were confirmed to respond to capsaicin at the end of each recording (if intraluminal infusion did not evoke action potential discharges, capsaicin would be added through the serosal side of the esophagus to the same recorded C fiber from the esophageal-vagal preparation).

**Esophageal intraluminal acid infusion activates esophageal nodose C fibers after in vivo OVA inhalation.** To determine the effect of acid on esophageal vagal nodose C fibers, we performed extracellular single-unit recordings from nodose ganglia neurons with intact nerve endings in the esophagus using ex vivo esophageal-vagal preparations. Esophageal distension-evoked action potentials were compared before and after intraluminal acid infusion. According to our previous studies (29, 32), we defined the response to a particular stimulus as positive when the stimulus-evoked action potential discharge had a peak frequency of at least 2–3 Hz above the baseline activity. In esophageal nodose C fibers from naive animals, acid infusion (HCl in KBS, pH 2.1) for 30–120 min did not evoke action potential discharges (baseline vs. acid: 2 ± 0.26 vs. 2.4 ± 0.16 Hz, n = 10, P > 0.05). After washing out acid with fresh KBS (pH = 7.4) for 30 min, intraluminal infusion with capsaicin (1 µM) for 30–60 min did not evoke action potential discharges in these C fibers (baseline vs. capsaicin: 2 ± 0.26 vs. 2 ± 0.45 Hz, n = 10; Fig. 1C). In contrast, these C fibers were activated by capsaicin perfusion from the serosal side of the esophagus at the end of each recording (baseline vs. capsaicin: 2.0 ± 0.26 vs. 14.6 ± 2.7 Hz, n = 10, P < 0.05).

In OVA-sensitized animals, OVA inhalation in vivo was performed three times every 24 h, then esophageal-vagal preparations were used for extracellular recording to determine intraluminal acid infusion-induced effects on esophageal nodose C fibers. Strikingly, intraluminal infusion with acid (using HCl in KBS and adjusted pH = 2.1) significantly evoked action potential discharges in esophageal nodose C fibers (baseline vs. acid: 1.3 ± 0.18 vs. 4.7 ± 0.63 Hz, n = 8, P < 0.05). These activation responses started after acid infusion for about 20 min (19.6 ± 5.4 min, n = 8). After washing out acid with fresh KBS (pH = 7.4) for 30 min, intraluminal infusion with capsaicin (1 µM) for 20–30 min also evoked action potential discharges in these C fibers (baseline vs. capsaicin: 1.3 ± 0.18 vs. 4.5 ± 0.8 Hz, n = 8, P < 0.05; Fig. 1A, B, and D). These activation responses started after infusion with capsaicin for about 21 min (20.6 ± 4.5 min, n = 8). These data

![Fig. 1. Intraluminal acid activates esophageal nodose C fibers after mast cell activation by in vivo ovalbumin (OVA) inhalation. A: extracellular single-unit recording from ex vivo esophageal-vagal preparation for intraluminal acid infusion. B: typical traces of action potential (AP) discharges evoked by intraluminal acid infusion in esophageal nodose C fibers before and after esophageal mast cell activation. C: in naive animals, intraluminal acid or capsaicin infusion did not evoke action potential discharges in esophageal nodose C fibers (baseline vs. acid, or vs. capsaicin = 2 ± 0.26 vs. 2.4 ± 0.16 Hz, or vs. 2 ± 0.45 Hz, both P > 0.05, n = 10). D: in OVA-inhaled animals, intraluminal acid or capsaicin infusion significantly evoked action potential discharges in esophageal nodose C fibers (baseline vs. acid, or vs. capsaicin = 1.3 ± 0.18 vs. 4.7 ± 0.63 Hz, or vs. 4.5 ± 0.8 Hz, n = 8, both *P < 0.05). GP, guinea pig.

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demonstrated that intraluminal acid did not activate esophageal nodose C fibers in naive animals but consistently activated nodose C fibers in the esophagus pretreated with OVA inhalations in OVA-sensitized animals. Our previous study (31) demonstrated that the mast cell is one of the most important immune cells involved in antigen induction of acute allergic reactions in the esophagus, and our present data from this model may suggest that the antigen induction of acute allergic response could play an important role in acid-induced activation of esophageal nodose C fibers.

Esophageal intraluminal acid infusion activates nodose C fibers after in vitro esophageal tissue challenge with OVA. To specify the role of mast cells in acid-induced activation of esophageal nodose C fibers, we induced mast cell activation by in vitro tissue antigen challenge and determined whether mast cell activation allowed acid infusion to activate esophageal nodose C fibers.

Mast cell activation by in vitro OVA tissue challenge was performed according to our previous study (30). We first performed extracellular single-unit recordings on nodose C fibers. This was followed by intraluminal acid infusion for 30–90 min, and the responses of esophageal nodose C fibers were determined. This time, we observed that intraluminal infusion with acid (HCl in KBS with pH = 2.1) also significantly evoked action potential discharges in these nodose C fibers. These data supported a specific contribution of mast cell activation to intraluminal acid-induced activation of esophageal nodose C fibers.

Intraluminal infusion with inflammatory mediators does not activate esophageal nodose C fibers. Previous studies reported that intraluminal acid could induce ATP release from esophageal epithelium and cause an inflammation response to produce a variety of inflammatory mediators (7, 17). It is well known that these mediators are able to activate and/or sensitize sensory afferents in the gastrointestinal tract (3, 4). To rule out that acid-induced activation of esophageal nodose C fibers is secondary for the release of these mediators, we tested the hypothesis that intraluminal infusion with these mediators themselves did not evoke action potential discharges in esophageal nodose C fibers.

We performed extracellular single-unit recordings on nodose C fibers in ex vivo esophageal-vagal preparations from naive animals. Intraluminal infusion with ATP (10–100 μM) for 30–60 min did not evoke action potential discharges in esophageal nodose C fibers (baseline vs. ATP: 1.3 ± 0.3 vs. 1.7 ± 0.3 Hz, n = 11). After ATP was washed out with fresh KBS (pH = 7.4) for 30 min, intraluminal infusion with capsaicin (1 μM) for 30–60 min did not evoke action potential discharges in these C fibers (baseline vs. capsaicin: 1.3 ± 0.3 vs. 2 ± 0.5 Hz, n = 11; Fig. 3A). In contrast, when perfused from the serosal side of the esophagus, ATP mildly increased action potential discharges in these esophageal nodose C fibers.
potential discharges of the same C fibers (baseline vs. ATP: 1.3 ± 0.3 vs. 2.9 ± 0.4 Hz, n = 11, P < 0.05). Similarly, capsaicin activated these C fibers when perfused through the serosal side (baseline vs. capsaicin: 1.3 ± 0.3 vs. 7.5 ± 1.5 Hz, n = 11, P < 0.05).

Intraluminal infusion with a previously reported “inflammation soup” (IS; including bradykinin, PGE2, serotonin, and histamine, all at 10 mmol/l) (4) for 30–60 min did not evoke action potential discharges in esophageal nodose C fibers (baseline vs. IS: 2.1 ± 0.4 vs. 2.3 ± 0.3 Hz, n = 15). After IS was washed out with fresh KBS (pH = 7.4) for 30 min, intraluminal infusion with capsaicin (1 μM) for 30–60 min did not evoke action potential discharges in these C fibers (baseline vs. capsaicin: 2.1 ± 0.4 vs. 2.4 ± 0.4 Hz, n = 15). When perfused from the serosal side of the esophagus in 8 of these 15 C fibers, IS increased action potential discharges among those C fibers (baseline vs. IS: 1.0 ± 0.31 vs. 5.8 ± 0.8 Hz, n = 8, P < 0.05). Similarly, capsaicin activated these C fibers when perfused through the serosal side (baseline vs. capsaicin: 1.0 ± 0.31 vs. 12.8 ± 1.65 Hz, n = 8, P < 0.05).

These data indicated that intraluminal ATP and inflammatory mediators were not able to activate esophageal nodose C fibers, probably due to the resistance of the epithelial barrier, which prevented these mediators to reach the nerve terminals.

Mast cell activation induces impairment of esophageal epithelial barrier function. Under the light microscope, we examined esophageal specimens (cross-sections with H-E staining) from animals that received treatments of KBS (pH = 7.4; Control group), ex vivo intraluminal acid (HCl in KBS, pH = 2.1) infusion (Acid group), and OVA inhalation in vivo plus ex vivo intraluminal acid infusion (OVA + Acid group). No severe injuries, including ulceration and erosion, were observed among these three groups. Based on reported methods (12), the inflammation grades were calculated from combined grades of inflammatory cells (mainly neutrophils) in the epithelial layer, basal cell hyperplasia, elongation of lamina propria papillae into the epithelium, and eosinophil infiltration. The esophageal inflammation grades mildly increased in animals treated with either acid infusion (5.78 ± 0.3) or OVA inhalation plus acid infusion (4.78 ± 0.3) compared to that from control (3.89 ± 0.9, vs. control, both P > 0.05, n = 4 in each group; Fig. 3, A–D). In contrast, increased esophageal mast cell infiltration was observed in both mucosa and muscle layers after antigen challenge in antigen-sensitized guinea pigs. Compared with those from naive animals (n = 3), OVA inhalation significantly increased mast cell numbers in the esophagus from OVA-sensitized animals (n = 4; naive vs. OVA inhalation: 8.33 ± 0.62 vs. 22.75 ± 2.07/cross-section, *P < 0.01).

Following extracellular recordings from esophageal-vagal preparations, esophageal mucosal epithelia were carefully dissected and TEERs were measured. In naive animals, acid infusion slightly decreased TEERs of esophageal epithelia (control vs. acid: 110 ± 7 vs. 92 ± 20 Ω·cm², n = 6–9, P > 0.05). In animals receiving OVA inhalation plus acid infusion, the TEERs of esophageal epithelia were significantly decreased (control vs. OVA + Acid: 110 ± 7 vs. 70 ± 14 Ω·cm², n = 6–9, P < 0.05; Fig. 4).

We also examined the expressions of tight junction proteins from these esophageal tissues. Compared to those from naive...
animals, both ZO-1 and occludin expressions were reduced in esophageal epithelia in animals receiving intraluminal acid infusion (ex vivo preparations) after in vivo OVA inhalations from OVA-sensitized animals. The ratio ZO-1/GAPDH significantly reduced from 91 ± 24% (naive, n = 4) to 17 ± 5% (OVA + Acid; n = 6, P = 0.027). The ratio occludin/NADPH reduced from 106 ± 32 (naive, n = 4) to 32 ± 11% (OVA + Acid; n = 6, P = 0.052; Fig. 5, A and B).

These data suggested that OVA inhalation plus ex vivo intraluminal acid infusion did not induce gross esophageal epithelial damage and severe inflammation, but led to reduced expression of tight junction proteins such as ZO-1 and occludin, which might contribute to impaired epithelial barrier function.

**DISCUSSION**

Clinical evidence supports a strong causal relationship between acid reflux and esophageal painful sensations such as heartburn and noncardiac chest pain, which can be successfully relieved by acid inhibition therapy (15). But direct evidence of acid-induced activation on esophageal nociceptive afferent nerve terminals is still lacking. Combining our established models for extracellular single-unit recording in esophageal-vagal preparations and mast cell activation by antigen tissue challenge or antigen inhalation in vivo, the present study for the first time demonstrated that intraluminal acid infusion activates nodose C fibers when esophageal mast cell activation is induced by either antigen inhalation or antigen tissue challenge in antigen-sensitized animals. Intraluminal acid infusion in the esophagus from antigen-inhaled animals did not induce severe epithelial damage but reduced the expression of tight junction proteins and increased epithelial permeability. These changes may allow intraluminal acid to reach sensory nerve terminals underneath and evoke action potential discharges in nociceptive C fibers.

Sensory nerves detect polymodal sensory stimuli through specific groups of ion channels expressed in neuronal cell bodies and nerve terminals. This enables them to transduct different stimulations in the peripheral tissues. Recently, several groups of ion channels have been proved to be involved in sensory tissue acidity. These mainly include acid-sensing ion channels, TRPV1, ionotropic purinoceptors (P2X), several Na⁺, K⁺, and Ca²⁺ channels, and proton-sensing G-protein-coupled receptors (11). At present, the information on acid transduction has largely been extrapolated from the studies carried out at the level of the cell bodies of isolated neurons. These studies have provided important information on acid sensing, but have given little attention to nerve phenotype or acid delivery kinetics—two variables that are critical in acid-induced pain. In the gastrointestinal tract, sensory afferents sensing acid can be processed by different ion channels, depending on the acidity of the local tissue environment, which can be varied in different segments of the gut. But acid transduction in esophageal sensory nerve terminals is still less clear. To better understand the acid-sensing mechanism in the esophagus, we have developed an ex vivo esophageal-vagal preparation to study vagal afferent functions in normal and inflamed esophagus (29, 32). Using these approaches, in the present study we demonstrated that intraluminal infusion with either acid or capsaicin in normal esophagus does not activate esophageal nodose C fibers, and the same C fibers actually can be activated by both acid and capsaicin if they can reach the nerve terminals by delivering from the serosal side of the esophagus.

Fig. 5. Expressions of tight junction proteins in esophageal epithelium: A: expressions of zona occludens-1 (ZO-1) and occludin in esophageal epithelium by Western blot. B and C: compared with controls (naive animals), intraluminal acid infusion in animals pretreated with OVA inhalation reduced the expression of ZO-1 (control vs. OVA + acid: 91 ± 24 vs. 17 ± 5% of GAPDH, P = 0.027, n = 4 in control and n = 6 in OVA + acid groups) and occludin (control vs. OVA + acid: 106 ± 32 vs. 32 ± 11% of GAPDH, P = 0.052, n = 4 in control and n = 6 in OVA + acid groups) in esophageal epithelium.

**Fig. 4. Transepithelial electrical resistances (TEERs) of esophageal epithelium. In controls (naive animals), intraluminal acid infusion decreased TEERs of esophageal epithelium (control vs. acid: 110 ± 7 vs. 92 ± 20 Ω·cm², P > 0.05, n = 6 in control and n = 9 in acid groups), and this decrease reached significance from animals pretreated with OVA inhalation (control vs. OVA + acid: 110 ± 7 vs. 70 ± 4 Ω·cm², *P < 0.05, n = 6 in control and n = 6 in OVA + acid groups).**
esophagus. This suggested that normal guinea pig esophageal epithelium might not be permeable to intraluminal acid infusion. This is in agreement with the observation from a previous study on mouse gastrointestinal vagal afferent fibers, using extracellular single fiber recordings in vitro, that application of HCl to the mucosal surface directly above the afferent receptive field did not activate either esophageal mucosal or tension receptors (20). But another study from the same group, using in vitro ferret esophageal-vagal preparations, showed that less than 2% of vagal afferents responded to high concentrations of HCl (19). The other group using in vivo single fiber recordings on rat esophageal vagal afferents also demonstrated that about 30% vagal afferent fibers responded to intraesophageal acid infusion (21). We thought that several factors might contribute to these different observations, such as, the species of animal, the afferent subtypes, the concentration and method of acid application, and the recording systems. In the present study, we did not study acid-evoked effects in esophageal nodose A6 fibers, which do not respond to noxious stimulation such as capsaicin (32) but might be activated by acid, as shown in airway nodose A6 fibers (14).

Previous studies have shown that esophageal acid infusion led to tissue damage and inflammation (7, 16, 25, 28). These experiments were carried out in acute or chronic acid-infusion models either in vivo or in vitro. In the present study, intraluminal acid infusion was performed in ex vivo esophageal-vagal preparations. This minimized the effects from those mediators released in the tissue when studying the acid-sensing process in the nerve terminals. Moreover, our data from the present study show that intraluminal infusion with these mediators did not evoke action potential discharges in esophageal nodose C fibers. Thus the observed activation effect by intraluminal acid in esophageal nodose C fibers is more likely a direct response in the nerve ending induced by acid itself.

The mast cell is an important resident immune cell in the tissues. It plays an important role in initiating allergic/inflammation reactions. Mast cell activation-released mediators regulate not only sensory nerves but also epithelial barrier functions in the proximity (2). It is well known that mast cell activation increases intestinal permeability (22, 24), which may involve mast cell chymase-dependent damage on intestinal epithelial tight junction proteins (9, 10). Tight junction proteins, such as ZO-1 and occludin, have been reported to express in human esophageal epithelium (13), and may play important roles in epithelial barrier functions. Our previous study showed that esophageal mast cells contain chymase in their granule content, which can be released after antigen-induced mast cell activation (30). Whether esophageal mast cell activation affects esophageal epithelial barrier function is still unknown. The data from the present study provided the first evidence that esophageal mast cell activation by antigen inhalation reduced the expressions of tight junction proteins and increased epithelial permeability to intraluminal acid. After in vivo antigen inhalation, intraluminal acid infusion subsequently activated esophageal nodose C fibers underneath. We further proved the specific involvement of mast cells by using a more specific tissue antigen challenge to induce esophageal mast cell activation, a well-defined model in our previous studies (29, 30), and demonstrated that intraluminal acid infusion also activated esophageal nodose C fibers afterward. The present results that acid activates esophageal nodose C fibers only after mast cell activation and that mast cell activation disrupts the esophageal epithelial barrier are novel findings. These provide us a model to further clarify acid-sensing mechanisms in esophageal vagal afferent subtypes.

We have reported that guinea pig esophagus is innervated by distinct subtypes of vagal nociceptive C fibers: the vagal jugular C fibers that are developmentally derived from the neural crest, and the vagal nodose C fibers derived from placodes (32). In the present study we only studied nodose C fiber subtypes. This is based on several considerations: first, esophageal nodose C fibers express more acid-sensitive ion channels/receptors (such as P2X) than jugular C fibers (33); second, the response of nodose C fibers to mast cell mediators in our ex vivo esophageal-vagal model is more well defined than in jugular C fibers (29); third, a previous study on acid-induced responses in guinea pig airway afferent subtypes demonstrated that citric acid application in receptive fields more frequently induced activation responses in nodose C fibers than in jugular C fibers (14). However, the manner in which neural crest vagal jugular C fibers respond to intraluminal acid infusion deserves further exploration. In the present study, intraluminal acid-infusion-evoked activation of esophageal nodose C fibers only occurred after esophageal mast cell activation. One critical question may arise: whether mast cell activation-induced sensitization of esophageal nodose C fibers also contributes to acid-induced activation of these C fibers. Based on the present data, we cannot rule out this possibility. This is a very important but complicated question, because mast cell-induced sensitization of esophageal nodose C fibers may involve several ion channels or receptors downstream, and the thresholds of these ion channels or receptors to acid stimulation are different. Based on the present model and results, we will continue to elucidate the contribution of peripheral afferent sensitization to acid sensing in the esophagus in a separate study.

In summary, the present study demonstrated that intraluminal acid infusion does not activate esophageal nodose C fibers from normal esophagus, but evoke profound activation responses after mast cell activation. Esophageal mast cell activation by antigen inhalation does not cause severe tissue damage and inflammation but increases epithelial permeability, which leads to intraluminal acid that is permeable to sensory nerve terminals underneath. These data not only provide for the first time direct evidence of acid-induced activation of nociceptive sensory nerve endings in the guinea pig esophagus but also establish a validated model to study acid-sensing processes in the esophagus. This finding may help to better understand the development of esophageal painful sensations (such as heartburn) in patients with nonerosive reflux disease, functional heartburn, or gastroesophageal reflux disease with asthma.

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DISCLOSURES

No conflicts of interest, financial or otherwise, are declared by the author(s).
AUTHOR CONTRIBUTIONS
Author contributions: S.Z., A.H., and S.Y. performed experiments; S.Z., Z.L., A.H., and S.Y. analyzed data; S.Z., Z.L., and S.Y. interpreted results of experiments; S.Z., Z.L., A.H., and S.Y. prepared figures; C.O. and S.Y. conceived and designed of research; S.Y. drafted manuscript; S.Y. edited and revised manuscript; S.Y. approved final version of manuscript.

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