Direct measurement of acid permeation into rat oesophagus

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Background and aims: The early responses of the oesophageal mucosa to acid perfusion may predict subsequent pathology. Mucosal responses to luminal acid may result either from acid permeating through the mucosa or from other unknown transduction mechanisms. In order to better understand the dynamics of acid permeation into the oesophageal mucosa, we measured interstitial pH (pHint) of the oesophageal basal epithelial layer, pre-epithelial layer thickness, and blood flow in rats in vivo during luminal acid challenge. A novel confocal microscopic technique was used in vitro to measure pHint from defined cellular sites in response to luminal and basolateral acidification.

Methods: 5-(and-6)-Carboxyfluorescein (CF) and carboxy-seminapthorhodofluor-1 (SNARF-1) fluorescence was used to measure pHi, by conventional and confocal microscopy, respectively, in urethane anaesthetised rats. Pre-epithelial layer thickness was measured optically with carbon particles as markers. Blood flow was measured with laser Doppler flowmetry.

Results: Luminal acidification failed to alter pHi in vivo and in vitro, but pHi was lowered by modest serosal acidification. Pre-epithelial layer thickness and blood flow increased significantly during luminal surface acid perfusion. Indomethacin had no effect on any acid related response.

Conclusion: In this first dynamic measurement of oesophageal acid permeation and pre-epithelial layer thickness, pHint was preserved in spite of high luminal acidity by two complementary techniques. Despite the apparent permeability barrier to acid permeation, oesophageal blood flow and thickness responded to luminal acid perfusion.

Acid related oesophageal disorders are currently receiving considerable attention as a consequence of their potentially severe morbilities. Although the late stages of acid related disease, including erosive oesophagitis and metaplastic transformation, are well described, the early responses to acid exposure, which must necessarily precede subsequent pathology, have been subject to lesser number of studies. Hence a grasp of these early responses could be of use in the prevention of oesophageal damage.

The oesophageal mucosa is highly resistant to refluxed concentrated gastric acid. Oesophageal resistance to acid is thought to reside in luminal pH gradients that may result from bicarbonate secretion in some species, resistance to acid permeation due to high resistance intercellular structures, and intrinsic cellular resistance due to acid-base transporters in the basal cell plasma membrane. As is the case with the gastric and duodenal mucosa, it is likely that unique defence mechanisms are in place that are regulated according to luminal pH. Unlike other acid exposed organs, however, the replicating cells within the basal pre-epithelial layer are 25–30 cell layers removed from the luminal surface. A question thus remains as to how far acid penetrates into the mucosa in the absence of injury. The luminal acid signal is rapidly transduced to submucosal structures, as acute luminal acid exposure produces responses such as changes in ion secretion, blood flow, and prostaglandin production, and chronic exposure to excess luminal acid arguably produces proliferative changes in the basal epithelial layer. It is not known however whether the signalling for these responses results from submucosal acidification or from alternate mechanisms. The observation that luminal acidification fails to produce much injury to the basal layer, whereas even modest acidification of the serosal surface has a far greater potential for producing cell necrosis, is consistent with minimal transmucosal acid permeation. Nevertheless, it is not known if these responses to luminal acid result from actual acid penetration into the basal layer or from indirect effects of luminal acid, such as from neural or other signalling mechanisms, as there are no published studies in which the depth of penetration of luminal acid into the mucosa was measured directly in vivo in the absence of injury.

To address the question of the depth of luminal acid penetration, and also to study the response of two putative mucosal defence mechanisms that may play important roles in the resistance of the stomach and duodenum to acid induced injury, we measured interstitial pH (pHint), pre-epithelial layer thickness, and blood flow in the stomach and duodenum of living rodents and a confocal microscopic technique used previously to measure extracellular pH in the stomach and colon. We hypothesised that the oesophagus serves as an effective barrier to penetration of luminal acid, preventing the submucosa from acidifying in the presence of luminal acid.

With in vivo microscopy and confocal in vitro microscopy, we demonstrated for the first time that rat oesophageal pHint was maintained despite high luminal acidity. Moreover, we also demonstrated that luminal acid can only superficially penetrate the oesophageal mucosa. Furthermore, our data are consistent with the involvement of mucosal blood flow and a possible participation of the pre-epithelial layer in mucosal protective mechanisms.

MATERIALS AND METHODS

Chemicals

5-(and-6)-Carboxyfluorescein (CF) and carboxy-seminalapthorhadofluor-1 (SNARF-1) were obtained from

Abbreviations: pHint interstitial pH, pHi, intracellular pH, CF, 5-(and-6)-carboxy fluorescein, SNARF-1, carboxy-seminalapthroughorhodofluor-1, FI, fluorescent intensity, BGI, background intensity.

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Molecular Probes Ltd (Eugene, Oregon, USA). N-2-hydroxyethylpiperazine-N’-2-ethanesulphonic acid (HEPES) and other chemicals were obtained from Sigma Chemical Co. (St Louis, Missouri, USA). Krebs solution contains (in mM): 136 NaCl, 2.6 KCl, 1.8 CaCl₂, and 10 HEPES at pH 7.0. Krebs solutions were titrated to pH 6.5, 7.0, or 7.4 with 1 N NaOH (2–5 mM added Na⁺), or titrated to pH 1.5 or 1.0 with 5 N HCl (15–25 added Cl⁻) Subsequent to pH titration, all solutions for in vivo studies were adjusted to 300 mosm with NaCl. For luminal acid perfusion, Krebs solution was pH 1.5 or 7.4 Krebs solutions. The basolateral chamber was perfused with pH 7.4 or pH 6.5 solutions containing 0.1 mM SNARF-1. SNARF-1 excitation was by the 488 nm line of an Ar laser, with emissions collected simultaneously at 580 (25) nm and 640 (30) nm, using two detectors. The fluorescence intensity ratio of 640 nm/580 nm was used to calculate pH values based on an experimentally established calibration curves. SNARF-1 was not included in perfusates with pH <6.5 due to lack of solubility at low pH. Intersitial pH was calculated from a calibration curve determined in the confocal microscope with 0.1 mM SNARF-1 in NaCl media of pH 6.0–8.0. A single point calibration was performed for each experiment to standardise pH calculation for daily instrument settings.

**Confocal microscopy**

**Tissue preparation**

All confocal microscopy of rat tissues was performed under the approval of the University of Indiana IUCAC. The technique was modified from that developed previously for measurement of extracellular pH in colonic crypts by confocal imaging. Male Sprague-Dawley rats weighing 250–300 g (Charles River) were sacrificed with halothane vapour, and the oesophagus was excised, flushed, and muscle stripped. The isolated mucosal sheets were kept in ice cold Dulbecco’s minimum Eagle’s medium (Invitrogen, Carlsbad, California, USA). All tissue was used within two hours of isolation. The muscle stripped mucosa was mounted in a microscope chamber, permitting independent control of luminal and basolateral perfusates. The chamber was custom manufactured (Summit Precision Machining, Baltimore, Maryland, USA) and is a modification of our microscopy chamber for study of polarised functions in cultured epithelia. Perfusates were run continuously throughout experiments at 1 ml/min at room temperature, through luminal and basolateral chambers having volumes of 25 µl and 15 µl, respectively. Perfusates for luminal perfusion were based on Krebs solution, titrated to pH 7.4 or pH 1.5. Basolateral perfusates were based on physiological saline containing in mM: 130 NaCl, 5 KCl, 2 CaCl₂, 1 MgSO₄·7H₂O, 25 mannose, 20 HEPES, and 0.1 mM carboxy SNARF-1, pH 7.4 or pH 6.5.

**Measurement of interstitial pH (pH_{L}) in vitro**

The method was adapted from previously published methodology developed for measuring extracellular pH in which SNARF-1 was validated for emission ratiometric measurement of pH in extracellular tissue spaces in gastrointestinal epithelia using confocal microscopy. We used a Zeiss LSM510 confocal microscope with a 40× C-Apo water immersion objective lens. The apical chamber was perfused with pH 1.5 or 7.4 Krebs solutions. The basolateral chamber was perfused with pH 7.4 or pH 6.5 solutions containing 0.1 mM SNARF-1. SNARF-1 excitation was by the 488 nm line of an Ar laser, with emissions collected simultaneously at 580 (25) nm and 640 (30) nm, using two detectors. The fluorescence intensity ratio of 640 nm/580 nm was used to calculate pH values based on an experimentally established calibration curves. SNARF-1 was not included in perfusates with pH <6.5 due to lack of solubility at low pH. Intersitial pH was calculated from a calibration curve determined in the confocal microscope with 0.1 mM SNARF-1 in NaCl media of pH 6.0–8.0. A single point calibration was performed for each experiment to standardise pH calculation for daily instrument settings.

**In vivo microscopy**

**Animal preparation**

All animal studies were conducted under protocols approved by the West Los Angeles VAMC IUCAC. In vivo microscopy was performed, as modified from prior studies of the stomach and duodenum and from prior studies of pH in mouse tumours. Male Sprague-Dawley rats weighing approximately 300 g (Harlan Laboratories, San Diego, California, USA) were fasted overnight but had free access to water. After urethane (1.25 g/kg) anaesthesia, the rat was placed supine on a plastic stage. Body temperature was maintained at 36–37°C by a heating pad and heat lamp. A tracheal cannula was inserted and saline was continuously infused through the left femoral vein at a rate of 1.08 ml/h using a Harvard infusion pump. Arterial blood pressure was monitored via a catheter placed in the left femoral artery. The abdomen was opened via a 3 cm midline incision and the anterior wall of the forestomach and pre-epithelial layer thickness using carbon particles. A light rod was used to illuminate the mucosal surface at an acute angle in order to visualise the surface detail of the mucosa. Arrow indicates carbon particles on the luminal epithelial layer surface. In (A), the focal plane is at the pre-epithelial layer surface, at the level of the carbon particles. In (B) The focal plane has been lowered to that of the mucosal surface. Note that the carbon particles cluster is out of focus whereas the mucosal surface is in focus. The Z axis travel of the microscope between the two focal planes is considered to be equivalent to the thickness of the overlying pre-epithelial layer.
lower oesophagus was incised using a miniature electrocautery. A concave stainless steel disk (16 mm diameter and 1.2 mm deep) with 3 mm central aperture was fixed watertight on the oesophageal mucosal surface with a silicone plastic adherent (Silly Putty; Binney and Smith Co., Easton, Pennsylvania, USA). A right angle laser Doppler probe was placed beneath the exposed oesophageal mucosa to measure oesophageal blood flow. A thin plastic coverslip was fixed to the disk with the silicone adherent to permit closed perfusion with solutions (total volume 50 ml; rate 0.25 ml/min) using a Harvard infusion pump. After correctly placing the chamber on the mucosa, a small area of mucosa was isolated from extraneous secretion, and exposed only to perfusate, by virtue of being surrounded by silicone putty and being covered with the perfused chamber. Rats were injected with 0.5 ml of CF 5 mg/kg intravenously in physiological saline five minutes prior to CF loading. We found that this autofluorescence was negligible compared with the high fluorescence of CF stained oesophageal mucosa excised from sacrificed rats. Mucosae were used as the focal plane. Images were captured every five minutes. Emission intensity was measured 10 seconds before and after each designated time point. The paired readings needed to calculate a fluorescence ratio were thus taken at a maximum of 20 seconds apart. Image analysis was performed on the recorded images: three small areas of oesophageal submucosa between microvessels were selected at random and then followed throughout the experiment. Fluorescence intensity of the selected area was measured using an 8 bit image analyser software (Image-Pro Plus v. 1.3; Media Cybernetics, Silver Spring, Maryland, USA). The fluorescence intensity at 495 nm divided by that at 450 nm and the resulting ratio was converted to pH by using an in vitro calibration curve according to the equation:

\[
\text{Fluorescence ratio} = \frac{(FI_{495} - BGI_{495})}{(FI_{450} - BGI_{450})}
\]

where \(FI\) = fluorescent intensity and \(BGI\) = background intensity.

BGI was defined as the intensity of oesophageal mucosa prior to CF loading. We found that this autofluorescence was homogeneous throughout the field and was close to 0 even with high camera gain and exposure time. BGI was therefore negligible compared with the high fluorescence of CF stained oesophageal mucosa. The mean of the ratios calculated from the three selected areas was defined as the ratio for that time period.

To demonstrate that in situ measurements were comparable with those measured in solution, in vitro tissue pH calibration was performed using conditions comparable with those in vivo experiments but with the use of muscle stripped oesophageal mucosa excised from sacrificed rats. Mucosae were equilibrated in CF free buffer solutions and were then placed in 5 \(\mu\)M CF containing solutions at the same pH for 30 seconds. The tissue was then examined by fluorescence microscopy. Figure 1 depicts a typical calibration curve of the fluorescence ratio versus pH for the tissue pH calibration. The tissue calibration curve was similar to in vitro calibration.

**Optical system**

For fluorescent microscopy, a modified multipurpose microscope (Carl Zeiss, Inc., Germany) with a 10x objective (NA 3.0; Rolyn Optics, Covina, California, USA) was used. The light source was a 100 W mercury lamp (Chiu Technical). A filter holder was placed in front of the lamp for manual replacement of 450 and 495 nm narrow bandpass excitation filters (Chroma Inc., Brattleboro, Vermont, USA). The television system consisted of a CCD colour camera (Optronics Engineering, Goleta, California, USA), a television monitor (Sony, Japan), and an S-VHS videotape recorder (Sony Electronics Inc., Japan). The CCD colour camera was connected to an Intel Pentium based IBM compatible microcomputer with FlashPoint framegrabbing videographic card (Integral Technologies, Inc., Silver Spring, Maryland, USA), and the captured image was digitised and stored on the hard disk of the computer for later analysis.

**Image analysis**

Fluorescent images of the microscopically observed chambered segment of oesophageal mucosa at 515 nm emission were captured and recorded using the computer hard disk drive. Capillary loops above the network of submucosal microvessels, which were assumed to be located in the subepithelial or basal epithelial layer of rat oesophagus, were used as the focal plane. Images were captured every five minutes. Emission intensity was measured 10 seconds before and after each designated time point. The paired readings needed to calculate a fluorescence ratio were thus taken at a maximum of 20 seconds apart. Image analysis was performed on the recorded images: three small areas of oesophageal submucosa between microvessels were selected at random and then followed throughout the experiment. Fluorescence intensity of the selected area was measured using an 8 bit image analyser software (Image-Pro Plus v. 1.3; Media Cybernetics, Silver Spring, Maryland, USA). The fluorescence intensity at 495 nm divided by that at 450 nm and the resulting ratio was converted to pH by using an in vitro calibration curve according to the equation:

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curves constructed using aqueous solutions containing 5 µM CF in Krebs solutions at varying pH, as described previously.\(^{16,17}\)

Measurement of pre-epithelial layer thickness
Pre-epithelial layer thickness was measured using a modification of previously described techniques.\(^{25,26}\) Graphite particles were placed over the mucosa to delineate the luminal surface of the pre-epithelial layer. The oesophageal mucosal surface was visualised with a fiberoptic light rod (American Optical Scientific Instruments, Buffalo, New York, USA) at 45° from the mucosa in order to provide reflectance images of the epithelial surface (fig 2). The microscope was alternately focused from the oesophageal surface to the graphite pre-epithelial layer. The distance of vertical travel of the microscope objective was measured by using a digital Z axis measuring device (Quick-Check, Metronics, Bedford, New Hampshire, USA) connected to the microscope, providing a measure of pre-epithelial layer thickness.

Measurement of oesophageal blood flow
We simultaneously measured oesophageal blood flow and pH\(_{e}\) using a technique modified from our prior measurements of blood flow and pH in gastric and duodenal epithelia.\(^{16,17}\) A right angle probe (#H41-3667; Vasomedics Inc., St. Paul, Minnesota, USA) was placed beneath the oesophageal surface. Blood flow was measured as the voltage output of the laser Doppler instrument (LaserFlo BPM803A; Vasomedics Inc.) and was expressed relative to basal measurements made during the last 4–5 minutes of the dye loading period.

Acid perfusion protocol
Oesophageal mucosae were superfused with pH 7.0 Krebs for 15 minutes, after which the perfusate was changed to pH 1.0 for 40 minutes. In some rats, indomethacin pretreatment (5 mg/kg subcutaneously) was given 30 minutes before acid perfusions.

Statistics
All data are expressed as mean (SEM). Comparisons between groups were made by one way analysis of variance (ANOVA) followed by Fisher’s least significant difference test. A p value of less than 0.05 was taken as significant.

RESULTS
In vitro measurements of oesophageal pH\(_{e}\)
We used confocal microscopy of dye loaded oesophagus in vitro to localise the fluorescent dye reporting pH\(_{e}\). SNARF-1 was chosen due to its pK\(_a\) and physical properties that resemble fluorescein dyes, and because of its dual emission characteristic that can be used with confocal microscopy.\(^{19,28}\) An in vitro preparation was used to facilitate perfusion of the basolateral surface. Figure 3 depicts fluorescence distribution in the confocal X-Z plane (optical cross section) using images of rat oesophageal mucosa from the same tissue location under several conditions: before dye perfusion (fig 3A) and subsequently after either luminal (fig 3B) or luminal plus serosal (fig 3C) addition of SNARF-1 dye and 10 minutes following removal of dye from both perfusates (fig 3D). Dye uptake occurred in an intensely fluorescent (white appearing) band when exposed to luminal SNARF, and had a more mottled appearance when SNARF was perfused over the basolateral surface. Figure 4 depicts fluorescence distribution in the confocal X-Z plane (optical cross section) using images of rat oesophageal mucosa from the same tissue location under several conditions: before dye perfusion (fig 3A) and subsequently after either luminal (fig 3B) or luminal plus serosal (fig 3C) addition of SNARF-1 dye and 10 minutes following removal of dye from both perfusates (fig 3D). Dye uptake occurred in an intensely fluorescent (white appearing) band when exposed to luminal SNARF, and had a more mottled appearance when SNARF was perfused over the basolateral surface of the preparation. Notably, SNARF was excluded from the portion of the epithelium farthest from the luminal or serosal surfaces (black band between fluorescent areas). As seen, dye introduced from the serosal perfusate permeated the intercellular space between the basal epithelial cells. To further localise the dye, serial X-Y (plane of the oesophageal mucosa) confocal sections were obtained in tissue 10 minutes after perfusion with SNARF-1 over the luminal and serosal mucosal surfaces (fig 4). A copy of fig 3C is included for orientation, with broken lines depicting the focal planes for each subsequent panel of fig 4. Note that SNARF-1 stained the flat polygonal cells of the stratum corneum (fig 4A), did not stain the middle epithelial layer (fig 4B).
4B), but when applied to the basolateral surface stained the basal cell layer and submucosa (fig 4C, 4D, respectively). These data are consistent with SNARF-1 penetration of the outermost 2–3 pre-epithelial layers of the stratum corneum and entering the flat polygonal cells of the outermost layer. The high SNARF-1 permeability of these outer cells suggests they have compromised viability, or at least compromised membrane integrity. In contrast, SNARF-1 staining at the basal layer appears to be exclusively extracellular, with dye permeating only the lateral spaces between basal cells (fig 4C, arrow). Staining of submucosal tissue in fig 4D suggests that SNARF may be adhering to certain proteins in the extracellular matrix, potentially explaining the retention of fluorescence after removal of SNARF-1 from the perfusates. Notably, dye was completely excluded from the remainder of the epithelium, which, by inference, is the stratum spinosum. After localising the dye to the basal pre-epithelial layer, we then imaged this pre-epithelial layer in the presence of acid perfused over the apical and basolateral mucosal surfaces. Figure 5 depicts SNARF-1 emission ratio images in the X-Z focal plane after perfusion of the basolateral surface with SNARF-1 for 10 minutes. Pseudocolour (green-acid; red-alkaline) corresponds to pHint. SNARF-1 could not be added to the apical perfusate as it precipitates at strongly acidic pH. Figure 5A, 5B, and 5D qualitatively demonstrate lack of change in pHint when luminal pH is changed from pH 7.4 to 1.5. This is quantitatively confirmed by the time course experiment shown in the bottom curves (A) and (B). Ability to elicit and detect changes in pHint in the basal layer is demonstrated by perfusion with a mildly acidic (pH 6.5) solution in the basolateral chamber (fig 5C and bottom curves (A) and (B)). These data are consistent with the lack of transmucosal acid permeation even in the presence of physiological transmucosal acid gradients.

**Measurements of oesophageal pHint in vivo**

In order to measure the depth of acid penetration into the mucosa in our in vivo model, we measured pHint using a technique in which CF (free acid), injected intravenously, leaked into the interstitial space. CF was used in preference to the structurally related SNARF-1 because our fluorescence microscope used for in vivo studies was equipped only for excitation ratio measurements, necessitating the use of a fluorophore appropriate for these measurements, such as CF. We continuously observed the mucosa under transillumination, injected CF intravenously, and then observed the development of fluorescence. As seen in fig 6, the submucosal vessels fluoresced shortly after CF injection, with transfer of fluorescence out of the vascular space occurring within 30 seconds after CF injection. Overall, CF fluorescence reached a maximum at 20 minutes, then gradually decreased, and remained at a level sufficient to obtain reproducible fluorescence ratios for 60 minutes, with a stable fluorescence intensity ratio (F495/F450) during perfusion with pH 7.0 buffer (fig 7). The ratio however decreased...
10 minutes after the rats were killed by intravenous air injection at 60 minutes. Basal pHₙₑₚ was measured during perfusion with pH 7.0 buffer perfusion, was 7.25 (0.05) (n=22). As shown in fig 8, rat oesophageal pHₙₑₚ was unchanged during perfusion of luminal applied pH 1.0 solution, further supporting our hypothesis that the oesophageal epithelium is an effective barrier to acid diffusion.

As prostaglandins enhance mucosal injury in experimental gastritis models in contrast with the gastroduodenal mucosa, we examined the effect of indomethacin pretreatment on pHₙₑₚ during perfusion with pH 7.0 or 1.0 solutions on oesophageal defence mechanisms in the absence of injury. Indomethacin had no effect on pHₙₑₚ in our system.

Measurement of pre-epithelial layer thickness

We then measured changes in relative pre-epithelial layer thickness to determine if it increased during luminal acid perfusion, suggestive but not conclusively implicating a protective role for the surface pre-epithelial layer. Pre-epithelial layer thickness varied from 0 to 67 µm, with a mean thickness of 29 (2) µm (n=67, 2-3 measurements/rat), and did not increase immediately in response to acid perfusion. Due to the delayed and gradual increase in pre-epithelial thickness, we were only able to demonstrate a statistically significant increase in pre-epithelial thickness after a 40 minute perfusion with pH...
overlying the oesophageal mucosa. The pre-epithelial layer was significantly thickened during pH 1 perfusion but was not affected by indomethacin (Indo) intraperitoneally. *p<0.05 versus pH 7 perfusion by ANOVA.

1.0 solution (p<0.05) compared with the thickness increase during perfusion with a pH 7.0 solution (fig 9).

Measurement of oesophageal blood flow

Hyperaemia during luminal acid challenge is another means by which gastrointestinal mucosa are protected during acid perfusion.\(^{32,33}\) We thus measured blood flow in the oesophagus using laser Doppler flowmetry. Figure 10 depicts relative oesophageal mucosal blood flow in rats perfused with either neutral or acidic solutions. In control rats perfused with neutral solutions, relative mucosal blood flow gradually decreased to 88 (5)% of baseline (n=7). During acid perfusion, mucosal blood flow significantly increased compared with control rats (p<0.05). Pretreatment with indomethacin had no effect on this increase in blood flow. This increased blood flow indicates that submucosal structures such as blood vessels can be influenced by luminal acid, in the absence of transepithelial acid diffusion, consistent with an indirect transduction mechanism. Increased blood flow and pre-epithelial layer thickness in response to acid is reminiscent of the gastroduodenal protective response to luminal acid.

**DISCUSSION**

We have demonstrated that intravenously injected water soluble pH indicating fluorescent dyes can permeate the interstitial space of the basal pre-epithelial layer of the rat oesophagus, serving as useful markers for acid penetration into the mucosa. We found that the pH\(_m\) of the basal layer of the rat oesophagus was unchanged when the luminal surface was exposed to high luminal acidity in vitro and in vivo, although pH\(_m\) decreased during exposure to modestly acidic solutions exposed to the basolateral surface of the preparation. We also found that pre-epithelial layer thickness and blood flow increased in response to luminal acid in rat oesophagus in vivo. We could not find any effects of indomethacin on our measurements.

The use of fluorophores to measure pH\(_m\) has been described in the literature for several decades. In our case, we used two dyes, both of which are highly water soluble and thus excluded from viable cells having an intact plasma membrane. These dyes, as shown by others, readily migrate from the vascular space or from a mucosal bathing solution to the intercellular interstitium, where they can faithfully record pH\(_m\). The emission light path of confocal microscopes is split, enabling simultaneous imaging of the same field at two wavelengths. Dyes such as SNARF-1 take advantage of this feature as they have a pH dependent measurable shift of emission wavelength. Conventional epifluorescent microscopes, such as used for the in vivo studies, do not have emission beam splitters. Hence fluorophores such as CF are used, in which the pH dependent ratio of fluorescence intensity at a single wavelength is measured while alternating between two excitation wavelengths. This is the more usual method used for FRIM based pH measurements (fluorescence ratio intensity measurement).

Measurements of pH\(_m\), pre-epithelial layer thickness, and blood flow all reflect the response of the epithelium to acid challenge. The 40–70 minute measurement period used for the in vivo studies is comparable with the duration of acute clinical reflux episodes\(^{15,34}\) although clinical disease is thought to develop after months or years of repeated acute reflux episodes. Through these types of measurements, we have been able to study mucosal physiology to understand which factors need to be regulated in order to prevent injury. In this fashion, these studies complement those designed with injury as an end point. We measured the pH\(_m\) of rat oesophagus with two novel but related methods: an in vivo microscopic technique and an in vitro confocal microscopic technique. Unexpectedly, changes in pH\(_m\) in the basal epithelial layer were not demonstrated, in spite of high luminal acidity. Basal interstitial pH decreased only when the blood supply was stopped by intravenous air injection in vivo or when the basolateral perfusate was changed from neutral to pH 6.5 solution in vitro. These findings indicated that the pH\(_m\) of the basal epithelial region was largely dependent on the pH of the blood or basolateral fluid rather than the pH of the lumen, and were consistent with low acid permeability of the outer epithelial layers to acid.

Oesophageal epithelial resistance to luminal acid has been extensively studied by Orlando et al., in addition to other investigators. The rabbit oesophageal epithelium is electrically tight (1000–2500 Ω×cm\(^2\)), with contributory structural components located predominantly in the stratum cornueum and upper stratum spinousum. Prolonged contact with luminal acid and pepsin alters the properties of the intercellular junctions so as to increase paracellular pathway permeability,\(^{15,34}\) enabling influx of protons into the intercellular space and subsequent mucosal acidification. Acidification of the intercellular space (that is, low pH\(_m\)) is important in that the oesophageal epithelial cell serosal membrane has a much greater susceptibility to acid injury than the apical membrane.\(^{15,34}\) Protons must have penetrated the mucosa to some degree as luminal
acid perfusion increased oesophageal blood flow. Furthermore, staining of the cytoplasm of the cells of the stratum corneum (Fig 4) indicates that there was acid penetration into the most superficial portion of the epithelium, where dye penetration into the cytoplasm indicated the presence of non-viable cells. Nonetheless, the degree of penetration, as judged from the in vivo and in vitro studies, was superficial as we were unable to acidify the basal stratum by exposing the luminal surface to pH 1.0, although it is possible that acid penetrated into the stratum spinosum where no dye was present to report pH. That no dye was present in the stratum spinosum also underscored the apparent impenetrability of this epithelial layer. These observations are thus consistent with intercellular dilution and buffering of back diffusing H$^+$ or, more likely, the impermeability of the stratum spinosum to aqueous solutes, including protons. Coupled with the increase in blood flow (and possibly of pre-epithelial layer thickness) in response to luminal acid perfusion, these data are consistent with mucosal responses to luminal acid being mediated by neural acid sensing mechanisms, as has been observed previously with the stomach and duodenum, or by non-neural pathways, rather than from direct penetration of acid through the epithelium into the basal pre-epithelial layer. These sensing mechanisms have been suspected on the basis of the ability of subjects with reflux disease to sense the presence of acid perfused into the oesophageal lumen, in addition to the aforementioned mucosal responses to luminal acid perfusion. Nevertheless, available immunohistochemical data have not been consistent with penetration of afferent nerves into epithelial strata more superficial than the basal cell layer. This raises the question as to how a luminal acid signal is transduced in the absence of proton permeation through the mucosa to the location of the afferent nerves. Although precausative factors from further influence on the pre-epithelial layer. In contrast with its effects on the stomach and duodenum, our data provide more support for this paradigm as indomethacin did not adversely affect measured values. In summary, the rat oesophageal epithelium is highly resistant to acidification from prolonged exposure to concentrated mineral acid presented to the apical surface. Despite this lack of acidification, unknown mechanisms transduce the presence of luminal acid into physiological responses such as increased blood flow and increased pre-epithelial layer thickness. It is likely that an impermeable structure combined with acid sensing neural mechanisms and a separate transmucosal signal transduction mechanism underlie the observed mucosal acid responses that occur despite the remarkable acid resistance of the rat oesophagus. Further understanding of these early acid responses could lead to new therapeutic targets for therapies designed to prevent the acid induced mucosal damage that underlies advanced oesophageal pathology.

ACKNOWLEDGEMENTS

We would like to thank Jonathan Lee for his technical assistance. Supported by Department of Veterans Affairs Merit Review funding, and NIH-NIDDK R01 DK54221, 5F30DK041301, and R01 DK54940.

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Gut 2003 52: 775-783
doi: 10.1136/gut.52.6.775

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