Research Article

Endoscopic Ex Vivo Evaluation of Bile Concentrations by Narrow Band Imaging: A Pilot Study

Roberta Maselli,^{1,2} Haruhiro Inoue,¹ Haruo Ikeda,¹ Manabu Onimaru,¹ Akira Yoshida,¹ Esperanza Grace Santi,¹ Hiroki Sato,^{1,3} Nikolas Eleftheriadis,¹ and Shin-ei Kudo¹

¹Digestive Disease Center, Showa University Northern Yokohama Hospital, 35-1 Chigasakichuo, Tsuzuki-ku, Yokohama 224-8503, Japan

²Surgical Sciences Department, Umberto I General Hospital, Sapienza University of Rome, Rome, Italy

³De La Salle University Medical Center, Dasmariñas, Philippines

Correspondence should be addressed to Roberta Maselli; roberta.maselli.md@gmail.com

Received 19 October 2014; Revised 16 February 2015; Accepted 23 March 2015

Academic Editor: Mala Banerjee

Copyright © 2015 Roberta Maselli et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Background. Bile juice plays a major role in duodenogastroesophageal reflux (DGERD). Several devices to directly measure the bile concentration have been proposed. We aimed to ex-vivo evaluate the bile concentration by narrow band imaging (NBI). **Method.** From six surgical cholecystectomies, the content of the gallbladders was aspirated and the total biliary acid (TBA) concentration was evaluated. 2 mL was employed for serial twofold dilutions. Each dilution was scoped. Images on white light (WL) and NBI were captured and grouped accordingly to NBI-appearance and TBA-concentration. **Results.** Nondiluted bile had a TBA-concentration of $61965 \pm 32989 \mu\text{mol/L}$. Final dilution (1:4096) had $1.16 \mu\text{mol/L}$. NBI and correspondent WL images were grouped into seven groups, and an NBI/Bile scale was created. **Conclusion.** The scale showed that not only NBI scale but also white light scale could be useful to predict the bile concentration. This initial study shows that NBI has a potential role in the detection of DGERD and further investigation is warranted to distinguish the presence and the concentration of bile, especially at very low TBA concentrations.

1. Introduction

In the pathogenesis of reflux esophagitis, acids, bile acids, and pepsin play a major role in damaging the esophageal epithelium [1–3]. However, the mechanisms by which the luminal contents affect the barrier function of the stratified epithelial cell layers remain unclear.

Duodenogastroesophageal reflux (DGER) of bile acids has been hypothesized to play an important role in the development of oesophageal symptoms, oesophagitis, Barrett's esophagus (BE), and oesophageal adenocarcinoma [4]. This is supported in animal models in which oesophageal bile exposure (with or without acid or pancreatic juice) by either direct perfusion or total gastrectomy with oesophagojejunal anastomosis results in severe oesophagitis, Barrett's metaplasia, and adenocarcinoma [5].

It has also been proposed that bile reflux into the airway contributes to extraoesophageal GERD manifestations,

although the evidence to support this contention is largely indirect.

The first report on narrow band imaging was published more 19 years ago [6] to demonstrate the contrast between the vascular pattern and the adjacent mucosa of the underside of the human tongue using five narrow band illuminations and three broadband illuminations.

The technology consists of placing narrow band pass filters in front of a conventional white light source to obtain tissue illumination at selected narrow wavelength bands. These bands produce the greatest contrast between vascular structures and the surrounding mucosa. Currently available NBI systems use 2 narrow band filters that provide tissue illumination in the blue (415 nm), corresponding to the main peak on the absorption spectrum of hemoglobin, and green (540 nm) spectrum of light, corresponding to a secondary hemoglobin absorption peak. NBI generates a darker field of view than its white light counterpart. Moreover, the presence

FIGURE 1: DGERD during esophagoscopy; note how severe the bile reflux appears on NBI (b) compared with correspondent white light image (a).

of bile and blood (e.g., after biopsy) obscures the view under NBI because these fluids strongly absorb the narrow band light [7]. The NBI red coloration of the yellowish organic fluids (bile, intestinal residues) is a common experience for endoscopists. More often, because of a normal low contrast between the bile and the pinkish esophageal mucosa, when switching to NBI the reddish bile reflux seems more severe than the one seen with white light (Figure 1). On the other hand, whether NBI is useful for DGER diagnosis and severity assessment has never been investigated. NBI therefore has a potential to allow a one-step, less invasive DGERD diagnostic tool.

The interaction of tissue structures with light, and consequently the appearance of the structures, is wavelength dependent. The different coloration of mucosal features with NBI is achieved by observation of light transmission at selected wavelengths (or colors).

Aim of this pilot study was to evaluate the utility of NBI to distinguish and categorize different concentration on bile in an in vitro model.

2. Materials and Methods

From six consecutive routinely performed surgical cholecystectomies, the entire content of the removed gallbladders was aspirated. Part of the bile juice was sent to the laboratory to evaluate the total biliary acid (TBA) concentration. TBA concentration was evaluated by a quantitative test (Modular P800, Hoffmann-La Roche Ltd., Basel, Switzerland); it was performed three times for each sample and finally expressed as $\mu\text{mol/L}$. Meanwhile, 2 mL of the bile was employed for serial twofold dilutions using sterile water as solvent. The dilutions achieved were then stored in white plastic test tube. Each test tube was finally scoped with the GIFH260 (Olympus Medical Systems Co. Tokyo, Japan), and images on white light and NBI were captured and stored in a dedicated software (SolemioENDO ProStudy; Olympus Optical Co Ltd., Tokyo, Japan). Stored images were then grouped accordingly to the NBI reddish appearance and TBA concentration.

Corresponding white light images were grouped accordingly. TBA concentration of each group was calculated and shown as mean $\pm$ standard deviation.

3. Results

Nondiluted bile from the gallbladders had a mean TBA concentration of $61965 \pm 32989 \mu\text{mol/L}$ (Group 1). It was possible to group NBI images into seven different groups according to the color intensity. A bile concentration scale was created, as showed in Figure 2. Sequential twofold dilutions had the following TBA concentration: Group 2 (1:4 dilution) had $17690 \mu\text{mol/L}$, Group 3 (1:16 dilution) had $3165.25 \mu\text{mol/L}$, Group 4 (1:64 dilution) had $412.31 \mu\text{mol/L}$, Group 5 (1:256 dilution) had $99.84 \mu\text{mol/L}$, and Group 6 (1:1024 dilution) had $29.88 \mu\text{mol/L}$. Final dilution (1:4096) had mean $1.16 \pm 1.31 \mu\text{mol/L}$ (Group 7).

4. Discussion

Bile acids can stimulate esophageal squamous epithelial cells and Barrett's epithelial cells to produce substances that might promote esophageal inflammation (e.g., IL-8 and COX-2) and bile acids can cause oxidative stress and DNA damage. Bile acids can induce esophageal squamous cells to change their gene expression patterns to resemble intestinal cells and can cause Barrett's epithelial cells to increase their expression of intestinal-type genes [8].

These data suggest that in patients with GERD the bile acids may contribute to the pathogenesis of symptoms, esophagitis, Barrett's metaplasia, and Barrett's related carcinogenesis [9].

Historically, measurement of DGER has been thought to be feasible by scintigraphy, endoscopic gastric aspiration techniques [10], ambulatory gastric pH monitoring [11], and more recently by endoluminal spectrophotometric techniques (Bilitec) and impedance monitoring. Bilitec 2000 is currently the only commercially available device that is proven effective in measuring bile reflux. Bilitec 2000 has

FIGURE 2: Bile concentration scale on white light (WL) and NBI. Each correspondent mean TBA concentration is reported on the right.

shown that patients with severe esophagitis [12] and patients with BE have more bilirubin reflux episodes than normal subjects and that these reflux episodes coincide with acid reflux episodes. Bilitec is an external probe to be inserted in the esophagus. In contrast to Bilitec, NBI is incorporated in the endoscope activated by a push of a button.

Considering the created scale, it seems that not only NBI scale but also white light scale could be useful to predict the bile concentration. However, this is an in vitro study and the contrast between the bile and the physiologic color of the mucosa must be taken into consideration and checked in the future. The bile concentrations were stored and scoped in a white plastic tube and this white background color could interfere with the NBI/white images. At the same time,

these tubes were chosen to have the same background, being impossible to reproduce the real in vivo conditions of normal gut.

Although future in vivo studies are needed to confirm this bile scale, this initial study shows that NBI has a potential role in the detection of DGERD and further investigation is warranted to distinguish the presence and the concentration of bile, especially at very low TBA concentrations.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

References

- [1] W. K. H. Kauer, J. H. Peters, T. R. DeMeester et al., "Composition and concentration of bile acid reflux into the esophagus of patients with gastroesophageal reflux disease," *Surgery*, vol. 122, no. 5, pp. 874–881, 1997.
- [2] D. C. Gotley, A. P. Morgan, and M. J. Cooper, "Bile acid concentrations in the refluxate of patients with reflux oesophagitis," *British Journal of Surgery*, vol. 75, no. 6, pp. 587–590, 1988.
- [3] G. Champion, J. E. Richter, M. F. Vaezi, S. Singh, and R. Alexander, "Duodenogastroesophageal reflux: relationship to pH and importance in Barrett's esophagus," *Gastroenterology*, vol. 107, no. 3, pp. 747–754, 1994.
- [4] F. Johnsson, B. Joelsson, C.-H. Floren, and A. Nilsson, "Bile salts in the esophagus of patients with esophagitis," *Scandinavian Journal of Gastroenterology*, vol. 23, no. 6, pp. 712–716, 1988.
- [5] M. F. Vaezi, S. Singh, and J. E. Richter, "Role of acid and duodenogastric reflux in esophageal mucosal injury: a review of animal and human studies," *Gastroenterology*, vol. 108, no. 6, pp. 1897–1907, 1995.
- [6] K. Gono, T. Obi, M. Yamaguchi et al., "Appearance of enhanced tissue features in narrow-band endoscopic imaging," *Journal of Biomedical Optics*, vol. 9, no. 3, pp. 568–577, 2004.
- [7] L. M. Song, D. G. Adler, J. D. Conway et al., "Narrow band imaging and multiband imaging," *Gastrointestinal Endoscopy*, vol. 67, no. 4, pp. 581–589, 2008.
- [8] W. K. H. Kauer, J. H. Peters, T. R. DeMeester et al., "Mixed reflux of gastric and duodenal juices is more harmful to the esophagus than gastric juice alone: the need for surgical therapy re-emphasized," *Annals of Surgery*, vol. 222, no. 4, pp. 525–533, 1995.
- [9] X. Chen, T. Oshima, J. Shan, H. Fukui, J. Watari, and H. Miwa, "Bile salts disrupt human esophageal squamous epithelial barrier function by modulating tight junction proteins," *American Journal of Physiology—Gastrointestinal and Liver Physiology*, vol. 303, no. 2, pp. G199–G208, 2012.
- [10] H. J. Stein, H. Feussner, W. Kauer et al., "Alkaline gastroesophageal reflux: assessment by ambulatory esophageal aspiration and pH monitoring," *The American Journal of Surgery*, vol. 167, no. 1, pp. 163–168, 1994.
- [11] S. Y. Ifrikhar, S. Ledingham, D. F. Evans et al., "Alkaline gastroesophageal reflux: dual probe pH monitoring," *Gut*, vol. 37, no. 4, pp. 465–470, 1995.
- [12] M. F. Vaezi, R. G. Lacamera, and J. E. Richter, "Validation studies of Bilitec 2000: an ambulatory duodenogastric reflux monitoring system," *The American Journal of Physiology*, vol. 267, no. 6, pp. G1050–G1057, 1994.