Mucin histochemistry of the columnar epithelium of the oesophagus (Barrett’s oesophagus): a prospective biopsy study

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SUMMARY Columnar epithelium-lined oesophagus (CELO) is an acquired disorder associated with a high incidence of cancer. CELO consists of three histological types of epithelium: gastric-fundic, junctional, and specialised columnar, the last resembling intestinal metaplasia of the stomach. In a previous study of CELO an incompletely differentiated variant of intestinal metaplasia secreting sulphomucins (type II B) was found. This was shown to be associated with well differentiated adenocarcinoma, as in the stomach. The purpose of this paper has been to define by histochemistry the mucin profile of CELO in 17 patients and to compare it with the mucin profile of the gastrooesophageal junction in 27 patients without CELO. In CELO a specialised columnar epithelium was always found and type II B intestinal metaplasia (with sulphomucins) showed the highest incidence (53%). In normal subjects, this type of intestinal metaplasia was found in only three of 27 cases. Type II B intestinal metaplasia has often been considered as a precancerous lesion or as an equivalent of dysplasia; consequently, its high incidence in our study on CELO raises the question of whether this lesion should be considered a high risk condition for adenocarcinoma of the lower oesophagus.

Columnar epithelium-lined oesophagus (CELO), or Barrett’s oesophagus, is a disorder in which the lower oesophagus is lined by columnar instead of squamous epithelium. CELO is usually an acquired condition complicating prolonged reflux oesophagitis1 and is of considerable interest because there have been several reports of a high incidence of adenocarcinoma arising in CELO, mostly in surgical specimens2 3 and in some biopsy material.4 5 Naef et al4 established the premalignant character of CELO, finding a 10% incidence of adenocarcinomas in a series of 140 biopsy samples. In another biopsy series, Savary et al5 found an even higher incidence (15.7%) of malignant changes in CELO.

Paul et al6 showed that CELO consisted of three histological types of epithelium: gastric-fundic, junctional, and specialised columnar. The last is a very distinctive epithelium,7 resembling intestinal metaplasia of the stomach. Jass8 showed that mucin histochemistry was helpful in further classifying specialised columnar epithelium. They found an incompletely differentiated variant of intestinal metaplasia secreting sulphomucins (type II B) associated with well differentiated adenocarcinoma in CELO, as previously described in the stomach by other authors.9 10 11

Using biopsy material we have attempted to define the mucin profile of CELO and of the normal gastrooesophageal junction by means of a prospective histochemical study carried out over two and a half years.

Material and methods

Forty four patients with symptoms of reflux oesophagitis were included in this study: 30 men and 14 women (mean age 52 years). Endoscopic examination and biopsy were performed on all these patients from January 1981 to June 1983. CELO was diagnosed in 17 patients (15 men and two women, mean age 51.5 years) on the basis of endoscopic changes and histological evidence of columnar lined epithelium. The other 27 patients (15 men and 12 women, mean age 52.3 years) did not show CELO by the above criteria. The number of biopsy
samples taken from each patient ranged from two to 13 (mean 4.76) in the 17 patients with CELO and from two to six (mean 3.55) in those without CELO.

All the samples were fixed in 3-6% neutral buffered formalin (modified Millonig's phosphate buffer) for 48 h and paraffin embedded. Serial sections, 4 μm thick, were cut from each sample and stained with the following: haematoxylin and eosin; alcian blue and periodic acid Schiff, pH 2-5, to differentiate acid from neutral mucins; and high iron diamine and alcian blue, pH 2-5, to differentiate sulphomucins from sialomucins.^13,14

**Results**

The histological and histochemical study of CELO showed heterogeneity. The incidence of the three histological types of epithelium of CELO as described by Paull et al^3^ is shown in Table 1. The two most common types were the junctional type (16) and the specialised columnar epithelium type (17). No dysplasia was found.

The histochemical reactions used for identifying epithelial mucins are shown in Table 2: neutral mucins stain with periodic acid Schiff, acid mucins are stained mainly with alcian blue, and sulphomucins are stained with high iron diamine. In CELO, both gastric fundic and junctional epithelium showed only neutral mucin secretion. Specialised columnar epithelium is characterised by intestinal metaplasia with goblet cells secreting acid mucins and intervening columnar cells (Fig. 1). Complete intestinal metaplasia type I is defined by the presence of columnar cells resembling normal enterocytes without mucus secretion. Incomplete intestinal metaplasia type II A is defined by the presence of columnar cells secreting neutral mucins or traces of sialomucins and type II B by the presence of columnar cells secreting sulphomucins (Fig. 2). Type II B showed the highest incidence and was fairly extensive but often focal and associated with other types of intestinal metaplasia. Table 3 shows the incidence of intestinal metaplasia subtypes in CELO. Forms of transition in terms of morphology and mucin profiles between these three well recognisable variants were also noted (I–II A, I–II A–II B, II A–II B). Nine of 17 cases of CELO (about half) showed the presence of sulphomucins. In 21 of 27 patients without CELO, biopsy samples

![Fig. 1 Incomplete intestinal metaplasia with goblet cells and intervening columnar cells. Haematoxylin and eosin. ×225](image)

*Recently Filipe et al^13^ proposed the term type II and type III intestinal metaplasia instead of type II A and type II B respectively.*

| Table 1 Incidence of the three histological types of epithelium in 17 cases of CELO |
|-----------------------------------------|-----------------|
| **Histological type** | **No of patients** |
| Gastric-fundic | 4 |
| Junctional | 16 |
| Specialised | 17 |
| *Overall total is greater than 17 patients because more than one type of epithelium occurred in some patients.* |

| Table 2 Histochemical techniques used to characterise gastrointestinal mucins |
|-----------------------------|--------------------------|
| **Subtypes** | **AB-PAS** | **HID-AB** |
| Neutral mucins | Magenta | O |
| Sialomucins | N-Acetyl | Purple-blue | Blue |
| | O-Acetyl | Blue | Blue |
| Sulphomucins | Blue | | Brown-black |
| O = negative staining. | PAS = periodic acid Schiff. | AB = alcian blue. | HID = high iron diamine. |

| Table 3 Incidence of intestinal metaplasia subtypes in CELO |
|-----------------------------------------|-----------------|
| **Intestinal metaplasia subtypes** | **No of patients** |
| I | 3 |
| I-IIA | 2 |
| II A | 3 |
| IIA–IIB | 3 |
| II B | 6 |
| Total | 17 |
| *Total of CELO with sulphomucins.* |
Recently, a mucin histochemical study of CELO suggested a similar histogenetic relation between type II B intestinal metaplasia and well differentiated adenocarcinoma of the oesophagus.

Our present prospective study emphasises the following aspects of CELO.

Nine of 17 patients with CELO showed type II B intestinal metaplasia (53%); this is in contrast with the conspicuous absence of sulphomucins in normal columnar epithelium of the gastrooesophageal junction. Our results are comparable with those of two recent histochemical studies of intestinal metaplasia, one in CELO and the other in gastric biopsy specimens. In a series of 10 cases of CELO (nine biopsy and one surgical specimen), Jass found type IIB intestinal metaplasia in three of 10 cases (focal only) and IIA intestinal metaplasia in five of 10 cases (usually extensive). In a prospective study of intestinal metaplasia in 1460 gastric biopsy specimens, Filipe et al showed a 8.7% incidence of type IIB intestinal metaplasia in 272 specimens with intestinal metaplasia. Our results as well as those of the above two studies show a higher incidence of type IIB intestinal metaplasia in CELO. Although the number of cases is small in our study, it would appear that the incidence of type IIB intestinal metaplasia is very high in CELO. With regard to a possible precancerous connotation of type II B intestinal metaplasia, our findings are consistent with the hypothesis that patients with CELO represent a very high risk cancer group.

Thompson et al suggested that Barrett's oesophagus (CELO) could be important as a precursor of adenocarcinoma arising in the region of the lower oesophagus and gastrooesophageal junction. In six biopsy specimens taken from the gastrooesophageal junction of patients with reflux oesophagitis but without CELO, intestinal metaplasia of either type II A (three) or type II B (three) was found. These results suggest that the same type of intestinal metaplasia could be present in patients with reflux oesophagitis, whether intestinal metaplasia is found above (that is, CELO) or at the level of an endoscopically normal gastrooesophageal junction. Consequently, the pathogenesis of adenocarcinoma of the lower oesophagus and of adenocarcinomas arising in the gastric cardia could be identical.

It would be of considerable interest to obtain a complete follow up of patients whose biopsy specimens contained II B intestinal metaplasia in order to see if type II B intestinal metaplasia could truly serve as a useful marker in reflux oesophagitis. Dysplasia has rarely been found in CELO. Among the biopsy specimens of 23 cases of CELO not associ-
ated with adenocarcinoma, Skinner et al^3 noted "low grade" (mild) dysplasia in only two cases. On the other hand, "high grade" or severe dysplasia often occurs adjacent to adenocarcinoma arising in CELO.4^16 When found in CELO, high grade or severe dysplasia is probably a good marker of malignant risk. Because of the low incidence of mild dysplasia in CELO, type IIB intestinal metaplasia could be a useful marker and regarded as a form of mild dysplasia in the screening for adenocarcinoma in CELO.

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References


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