A study to determine plasma antioxidant concentrations in patients with Barrett’s oesophagus

D M Clements, D A Oleesky, S C Smith, H Wheatley, D A Hullin, T J Havard, D J Bowrey

Background: Dietary questionnaire studies have suggested that patients with oesophageal adenocarcinoma are deficient in antioxidants. It is not known whether the same holds true for patients with the precursor lesion, Barrett’s oesophagus.

Aims: To evaluate the hypothesis that patients with Barrett’s oesophagus are deficient in antioxidants compared with patients without evidence of Barrett’s oesophagus.

Patients and methods: Plasma antioxidant profiles (copper, selenium, zinc; vitamins A, C, and E; carotenoids) were determined for patients with Barrett’s oesophagus (n = 36), patients with erosive oesophagitis (n = 32), and patient controls (n = 35).

Results: Patients with Barrett’s oesophagus had significantly lower plasma concentrations of selenium, vitamin C, β-cryptoxanthine, and xanthophyll compared with the other groups.

Conclusions: This study confirms the hypothesis that patients with Barrett’s oesophagus are deficient in certain antioxidants.

Sample size determination
In a previous study, we identified significant differences in the antioxidant concentrations between patients with chronic pancreatitis and healthy control subjects, and the magnitude of the differences was in excess of one standard deviation. For our current study, we assumed that the magnitude of the differences between Barrett’s and control patients would be more modest, in the order of 0.75 standard deviations. Assuming a significance concentration of 5% and a power of 80%, equal groups of 30 patients would be required.

Assay details
The biochemical parameters assessed were plasma concentrations of:

- the trace elements copper, selenium and zinc;
- vitamins A, C, and E;

A n epidemiological link between antioxidant deficiency and squamous cell carcinoma of the oesophagus has been established, most notably in China. Furthermore, antioxidant supplementation has been shown to produce modest reductions in the risk of progression of squamous dysplasia in this high risk population.

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The role of antioxidants in adenocarcinoma of the oesophagus is unclear. The principal risk factors for this cancer subtype are gastro-oesophageal reflux disease and increased body mass index. Ten studies have examined the diets of patients with oesophageal adenocarcinoma by means of structured questionnaires. Although the conclusions of these studies differed slightly, in general, patients were found to have lower intakes of antioxidants, citrus fruits, and raw vegetables compared with control populations. Two studies have examined selenium values in patients with Barrett’s oesophagus and concluded that high concentrations of this antioxidant were associated with a reduced risk of progression of Barrett’s oesophagus.

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No study to date has compared antioxidant concentrations in patients with Barrett’s oesophagus, the precursor lesion for adenocarcinoma, and lesser grades of oesophageal injury. Therefore we undertook our current study to evaluate the hypothesis that patients with Barrett’s oesophagus are deficient in antioxidants compared with patients with reflux who have no evidence of Barrett’s oesophagus.

Patients and methods
Plasma antioxidant profiles were determined for a prospectively enrolled cohort of subjects attending the endoscopy suite at the Royal Glamorgan Hospital, Llantrisant, UK, during the period May to September 2003. Our study was approved by the Bro-Taf local research ethics committee (protocol 03/5011) and the Royal Glamorgan Hospital research and development board. Each patient provided written informed consent for venepuncture.

Because antioxidants may act as reverse acute phase reactants, patients with occult inflammatory processes were excluded on the basis of either a raised C reactive protein concentration (> 10 mg/litre) or hypoalbuminaemia (< 32 g/litre). Additional exclusion criteria were the presence of gastroduodenal ulceration and a history of pancreatitis or previous foregut surgery, other than cholecystectomy.

Patients were divided into the following study groups:

1. 35 patients (33 men, three women) with Barrett’s oesophagus (defined by endoscopic columnar lined oesophagus and intestinal metaplasia on biopsy), mean age 57 years (range, 39–85);
2. 32 patients (20 men, 12 women) with Los Angeles grade B or C erosive oesophagitis, mean age 59 years (range, 35–77);
3. 35 patient controls (15 men, 20 women) free of reflux symptoms who had normal endoscopic appearances in the oesophagus, stomach, and duodenum, mean age 49 years (range, 20–72).

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the carotenoids (α carotene, β carotene, β cryptocyanxthine, lycopene, and xanthophyll).

Venous blood was collected from each subject after a six hour fast (immediately before endoscopy). Samples for trace element analysis were collected in trace element free sodium heparin Vacutainers (Becton Dickinson, Le Pont de Clais, France); those for vitamin analysis were taken into lithium heparin Vacutainers (Becton Dickinson) and transported to the laboratory in a light excluding container. All samples were centrifuged within 30 minutes of collection; separated plasma was divided into aliquots and stored in a −70°C freezer.

Samples were subsequently analysed in batches in the medical biochemistry department, University Hospital of Wales, Cardiff, UK. Copper and zinc were measured using flame atomic absorption on an FS-220 spectrophotometer (Varian Inc). Vitamins A, C, and E and selenium were measured by electrothermal graphite furnace atomic absorption on an FS-220 spectrophotometer (Varian Inc, Lexington, Massachusetts, USA). Selenium was measured using spectrophotometry with spectrophotometric detection.

Continuous data were compared using the unpaired *t* test, with significance assumed at the 5% level.

### RESULTS

Tables 1–3 summarise the antioxidant profiles. Patients with Barrett’s oesophagus had significantly lower plasma concentrations of selenium, vitamin C, β cryptocyanxthine, and xanthophyll compared with the other groups.

### DISCUSSION

It is unclear why some patients with gastro-oesophageal reflux disease develop Barrett’s oesophagus. Physiological oesophageal studies have shown that Barrett’s oesophagus is closely linked to poor lower oesophageal barrier function, hiatus hernia, and high levels of oesophageal acid (and bile) exposure. Nonetheless, even for similar oesophageal reflux exposure, end organ manifestations vary between individuals; genetic factors may be implicated. Our current study evaluated whether or not antioxidants could influence the development of Barrett’s oesophagus.

There are few publications on antioxidants in patients with Barrett’s oesophagus. However, several studies have evaluated antioxidant intakes in patients with oesophageal adenocarcinoma using dietary questionnaires. Antioxidant intakes in patients with reflux, with and without Barrett’s oesophagus. Antioxidants are

### Take home messages

- Patients with Barrett’s oesophagus were deficient in certain antioxidants—they had significantly lower plasma concentrations of selenium, vitamin C, β cryptocyanxthine, and xanthophyll than the other groups.
- It is possible that antioxidant supplementation could reduce the mortality risk for glandular dysplasia and adenocarcinoma in these patients and appropriate studies are needed to explore this issue.

### Table 1 Trace element concentrations

<table>
<thead>
<tr>
<th>Analyte</th>
<th>Barrett’s oesophagus (n = 36)</th>
<th>Erosive oesophagitis (n = 32)</th>
<th>Patient controls (n = 35)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Copper (μmol/l)</td>
<td>16.0 (15.0 to 16.9)</td>
<td>16.9 (15.5 to 18.3)</td>
<td>16.3 (14.9 to 17.7)</td>
</tr>
<tr>
<td>Selenium (μmol/l)</td>
<td>0.72* (0.67 to 0.78)</td>
<td>0.75 (0.71 to 0.79)</td>
<td>0.81 (0.74 to 0.87)</td>
</tr>
<tr>
<td>Zinc (μmol/l)</td>
<td>13.3 (11.4 to 15.1)</td>
<td>12.9 (12.2 to 13.5)</td>
<td>13.1 (11.6 to 14.6)</td>
</tr>
</tbody>
</table>

*Values shown are mean (95% confidence interval). *p* = 0.05 v patient controls.

### Table 2 Vitamin concentrations

<table>
<thead>
<tr>
<th>Analyte</th>
<th>Barrett’s oesophagus (n = 36)</th>
<th>Erosive oesophagitis (n = 32)</th>
<th>Patient controls (n = 35)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vitamin A (μmol/l)</td>
<td>2.17 (1.97 to 2.38)</td>
<td>2.01 (1.82 to 2.21)</td>
<td>1.97 (1.83 to 2.11)</td>
</tr>
<tr>
<td>Vitamin C (μmol/l)</td>
<td>18.3* (13.2 to 23.4)</td>
<td>26.6 (19.6 to 33.6)</td>
<td>27.1 (21.4 to 32.7)</td>
</tr>
<tr>
<td>Vitamin E (μmol/l)</td>
<td>26.7 (23.0 to 30.4)</td>
<td>29.1 (25.6 to 32.6)</td>
<td>28.3 (25.1 to 31.4)</td>
</tr>
</tbody>
</table>

*Values shown are mean (95% confidence interval). *p* = 0.02 v patient controls and *p* = 0.05 v erosive oesophagitis.
central to the cellular defence mechanism against oxidative damage—they catalyse the breakdown of hydrogen peroxide and fatty acyl lipid peroxides, in the presence of reduced glutathione, to water and the corresponding alcohols. Such peroxides are a source of potentially damaging free radicals, which can cause peroxidation of polyunsaturated fatty acids in the cell membrane. The toxic oxygen derivatives, which are neutralised by the antioxidants, are produced by normal cellular metabolic activity and by a variety of injurious agents.

Our main findings were that patients with Barrett’s oesophagus had significantly lower plasma concentrations of the antioxidants vitamin C, β cryptoxanthine, and xanthophyll compared with patient controls. What is the therapeutic potential of these observations? More information is available on squamous dysplasia and neoplasia, notably from China. The epidemiology of squamous carcinoma in high incidence regions of China (Linxian province) supports an association with antioxidant deficiency. Large scale antioxidant supplementation studies in this region revealed modest reductions in the mortality risk from oesophageal or gastric cardia malignancy. It is an appealing concept that the same will hold true for glandular dysplasia and adenocarcinoma.

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*J Clin Pathol* 2005 58: 490-492
doi: 10.1136/jcp.2004.023721

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