An unusual helical micro-organism found in the gut lumen of human subjects

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An earlier report described the discovery of a micro-organism in the form of a double helix in human small bowel biopsies. Mucosal biopsies of the stomach and small bowel obtained from patients with rheumatic diseases and dyspepsia by enteroscopy and gastroscopy were fixed for scanning electron microscopy to investigate the organism further. In 62% of biopsies, an organism in the form of a double helix with bifid ends, 5–30 μm long, was found lying free on the surface of the mucosa. The organism has been demonstrated in the stomach, duodenum and small bowel. Flagella were never seen to be associated with the organism. In spite of its helical form, the organism lacks many of the factors associated with spirochaete morphology. It is suggested that this, as yet unnamed organism, may be found throughout the length of the digestive tract. Its pathological significance is not known.

Introduction

A previous study demonstrated by scanning electron microscopy (SEM) that an organism in the form of a double helix, previously unknown, was present on the surface of the mucosa of human small bowel biopsies obtained at enteroscopy [1]. The organism, which appeared to be between 5 and 30 μm long, was invariably seen lying on, but never invading, the small bowel mucosa. It was thought to inhabit the true small bowel rather than the duodenum. Preliminary findings suggested that the organism was present in any individual in high numbers and, although often found in patients suffering from rheumatic complaints, it was subsequently found in patients investigated for non-ulcer dyspepsia who required endoscopic investigation of the upper gastrointestinal tract. This report describes further views of the organism, demonstrated by scanning electron microscopy (SEM), and suggests that it may be found not only in the true small bowel, but also in the stomach, duodenum and colon.

Materials and methods

Patients

The tissue biopsies examined in this study were taken from patients referred for endoscopic investigation of their gastrointestinal tract for various reasons; they were not chosen as part of a protocol. Those requiring enteroscopy [2] were suffering from forms of arthritis and were anaemic. The majority were taking non-steroidal anti-inflammatory drugs (NSAIDs) and were suspected to be bleeding from the small bowel [3]. Biopsies of small bowel mucosa were taken as part of the investigation of the patients’ anaemia, to identify lesions caused by NSAIDs and to demonstrate a normal villus pattern. Other patients requiring only gastroscopy and duodenoscopy were examined because of complaints suggestive of peptic ulcer disease. Gastric biopsies were taken to be examined for the presence of Helicobacter pylori and duodenal biopsies were taken to exclude villus atrophy. The patients were routinely prepared for endoscopic examination by being starved for at least 6 h before examination, and were sedated during the procedure with intravenous diazepam (Diazemuls; Dumex).

Collection and preparation of gastrointestinal biopsies

Enteroscopy was performed with an Olympus SIF-10 enteroscope (Olympus; Japan), with standard colonoscopy forceps. Mucosal biopsies of the small bowel and stomach were collected by a no-touch technique. A total of 53 biopsies was taken for examination. The specimens of mucosa were washed out of the forceps cups by agitation in 5 ml of ice-cold glutaraldehyde 2% in 0.05 M phosphate buffer, pH 7.4 (366 mOsms), and further processed within 48 h by the OTOTO method of
post-fixation [4–6]. They were dehydrated in a graded acetone series and critical point dried from liquid CO₂. Finally the specimens were sputter-coated lightly with gold and viewed in a JEOL JSM6310 SEM or a JEOL JSMT330 SEM (JEOL; Tokyo, Japan) operating at 15–20 KV.

The presence of *H. pylori* was shown by gastric mucosal biopsy. A single biopsy was tested by the direct urease test with the standard CLO test (TriMed, Carbon Court, Osborne Park, Western Australia). Each test was read after an interval of 30 min. No attempt was made to culture *H. pylori* from gastric biopsies to prove its presence.

**Results**

Examples of a double helical organism obtained from sites in the stomach, duodenum and small bowel are presented. Figs. 1–3 show examples of apparently identical organisms obtained from different sites of the gastrointestinal tract demonstrated by SEM. They were lying singly on the surface of the mucosa and in tangled groups occasionally associated with the orifices of mucus glands. They did not appear to invade the mucosal surface. The above examples are representative of the morphology of these helical organisms, which were seen to be of a regular shape and size for all the sites and various patients studied.

On an annual basis *H. pylori* is found in 18% of patients undergoing routine biopsy for upper gastrointestinal complaints in this unit (n > 1000). Of the 15 gastric biopsies investigated in the present study, five were found to be positive for the presence of *H. pylori*. The presence of the helical organism did not correlate with *H. pylori* status.

**Discussion**

A previous report [1] described double helical organisms obtained from the human small bowel by enteroscopy and described them as spirochaetes. However, on reflection, these organisms do not readily fulfil the criteria that define the primary characteristics of spirochaetes and may represent a separate group of organisms. Characteristically, the morphology of spirochaetes includes a single protoplasmic cylinder surrounded by a gram-negative cell wall. Axial fibrils and a complex of terminal periplasmic flagella comprising an axial filament lying within an outer membrane may be present.

The double helical organisms described here have none of these primary characteristics. They vary in length from c. 5 to 30 µm and are composed of a double helix, rather than a single spiral structure. The double helix appears to consist of two separate and identical spiral filaments, each filament terminating in a free fusiform end, giving the complete helix identical bifid ends. Flagella have never been seen to be associated with these organisms in any of the samples (>50). Furthermore, it is generally considered that true spirochaetes do not inhabit the small bowel.
[7, 8], but are commonly found in the mouth, large bowel and rectum – although one report refutes this [8].

The finding of the organism described here has been an incidental feature of the examination of biopsies taken for other reasons. No formal study of the frequency of these organisms in the gastrointestinal tract has been attempted, but it is evident from their common occurrence and the density at which they occur on mucosal biopsy specimens that they must represent frequent inhabitants of the stomach and small bowel. A review of previous studies of gastrointestinal organisms suggests that an identical organism has been demonstrated on the wall of the large bowel taken from a cadavar [9]. The results of the present study indicate that the presence of this helical organism is unrelated to a specific condition, i.e., rheumatoid arthritis or *H. pylori* status. However, the percentage of biopsies on which the organism was found in patients taking NSAIDs was 33%; in other cases it ranged from 75 to 87%. It must be
Fig. 3. Examples of helical organisms, obtained by enteroscopy from c. 60–100 cm distal to the pyloric sphincter in patients with rheumatoid arthritis and suspected of having associated lesions of the small bowel.

- **a.** A villus tip with terminal cleft; helical organisms lie scattered on the surface; mucus glands and individual enterocytes are apparent; bar = 10 μm.
- **b.** Helical organisms lying on the small bowel mucosa 100 cm distal to the pyloric sphincter; the outlines of individual enterocytes are visible; bar = 5 μm.

Emphasised that these findings are not statistically sound. It is concluded that this organism, of no known pathological potential, may be found in the human gastrointestinal tract, from stomach to colon.

References

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