Total gastrointestinal endoscopy in the management of Peutz-Jeghers syndrome

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Summary

Peutz-Jeghers syndrome was diagnosed in a 51-year-old woman presenting with iron deficiency anaemia. Upper gastrointestinal endoscopy and colonoscopy revealed several hamartomatous polyps in the stomach, duodenum and colon, which were removed. At a combined surgical-endoscopic procedure, 42 hamartomatous polyps were removed from the small intestine by snare polypectomy. This endoscopic procedure reduces symptoms, may protect against future intestinal obstructive episodes and their associated surgery, and may reduce the risk of developing gastrointestinal malignancy.

Keywords: Peutz-Jeghers syndrome; cancer; endoscopy

Peutz-Jeghers syndrome (PJS) is a rare autosomal dominant inherited disorder. It is characterised by hamartomatous polyps in the gastrointestinal tract and mucocutaneous melanin hyperpigmentation of the lips, mouth and less commonly the fingers and toes. The melanin spots tend to fade in adult life.1 The gene responsible for PJS has been mapped to chromosome 19p.2

The polyps in PJS may be multiple or solitary, and are most commonly present in the small intestine (64-96%), but also occur in the colon (60%) and in the stomach (24-49%).4 Patients typically present in the second or third decade of life with infaclature, ulceration and bleeding from polyps, resulting in anaemia, or with intestinal obstruction and intussusception.

Before the use of endoscopy, PJS was managed by laparotomy, manual reduction of intussusception, enterotomy, colotomy and surgical polypectomies or segmental resections.5 Repeated surgery, which is often necessary in PJS, may be complicated by the formation of peritoneal adhesions and short bowel syndrome.6 More recently, intraoperative endoscopy has been used to reduce symptoms from small intestinal polyps and reduce the frequency of emergency surgery.7

Between 2-13% of patients with PJS develop gastrointestinal cancer and a hamartoma-adenoma–carcinoma sequence has been suggested.8 Other malignant associations are bilateral breast cancer, pancreatic cancer, ovar-
intestinal obstruction in the future, it was decided to remove her small intestinal polyps endoscopically at laparotomy. At laparotomy, the surgeon could palpate approximately 20 polyps throughout the small intestine. A single enterotomy was made in the mid-jejunum at the site of a large polyp. An adult gastroscope, sterilised by 12 hours submersion in glutaraldehyde, was then passed into the small intestine. The endoscope was advanced in a retrograde direction to the duodenum and then antegrade to the terminal ileum with the surgeon helping to concertina the bowel over the gastroscope. A total of 42 polyps, the largest being 3 cm, were snared and removed by cautery. The surgeon assisted in retrieval of snared polyps by squeezing the detached polyps towards the enterotomy site. This saved time by eliminating the need to catch every detached polyp with endoscopic retrieval forceps and remove the gastroscope for each polyp. Total operative time was 90 minutes. Histological examination of the small intestinal polyps showed them all to be hamartomatous with no features of malignancy (figure 2).

There were no complications during or after the operation and the patient was discharged home after 7 days. Her anaemia responded to oral iron therapy.

Discussion

Patients with PJS have gastrointestinal polyposis, most commonly of the small intestine. They are often symptomatic, requiring surgery and there is an increased risk of developing gastrointestinal malignancy as well as other malignancies. Complete endoscopic examination of the gastrointestinal tract and polypectomy allows the patient with PJS a reduction in symptoms of anaemia, intussusception, and intestinal obstruction.

Patients with PJS under the age of 30 years often die of complications secondary to bleeding, intussusception and obstruction. However, patients over 30 years are more likely to die from gastrointestinal malignancy. There is some evidence for a hamartoma–adenoma–carcinoma sequence, although it has also been suggested that cancers can develop de novo from normal mucosa. The small intestine is the most common site for polyps in PJS, but malignancy in the small intestine is much less common than in the colon. An explanation for this may be that large polyps in the small intestine cause obstruction and are removed surgically before they become malignant.

Despite the development of long flexible endoscopes, complete endoscopic examination of the small intestine poses substantial technical difficulties, whether the endoscope is passed via the oral or anal route. At colonoscopy, only a short length of terminal ileum may be inspected. Using a 'push' enteroscope inserted orally, the proximal small intestine may be inspected but the ileum is often not reached. Sonde enteroscopy will more reliably reach the terminal ileum but has the disadvantages of being less controllable and without therapeutic capability. The combined surgical–endoscopic approach we used ensures a quick, thorough assessment of the entire small intestine. Small intestinal polyps not seen with barium imaging and not palpable through the bowel wall by the surgeon can be identified. Multiple polyps can be removed safely with the snare and retrieved for histology without the need for multiple enterotomies. However, the intra-operative surgical–endoscopic approach has the drawback of exposing the patient to the potential complications of general anaesthesia and laparotomy.

In addition to treating anaemia from chronic gastrointestinal blood loss, the total clearance of all polyps from our patient's gastrointestinal tract should provide some protection against the future development of gastrointestinal cancer and obstructive symptoms. Regarding future surveillance, some authors recommend performing gastroduodenoscopy, colonoscopy and barium imaging of the small intestine every two years to look for further polyp development. Breast and gynaecological screening have also been recommended in women. Although such surveillance and screening programmes may provide benefit and be justified, demonstrating efficacy of such programmes is difficult, given the rarity of PJS.

Summary points

- Peutz-Jeghers syndrome can be recognised by its cutaneous manifestations and these should be sought in patients presenting with anaemia, intestinal obstruction and intussusception
- the management of PJS includes total gastrointestinal endoscopy and polypectomy to protect against the development of gastrointestinal cancer and obstructive symptoms
- patients with PJS are also at risk of non-gastrointestinal malignancies and may benefit from screening
Abnormal liver function tests following inadvertent inhalation of volatile hydrocarbons

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Summary
The use of aerosols containing volatile hydrocarbons in conditions of poor ventilation can result in accidental over-exposure which can cause central nervous system effects and hepatic injury. We present a case in which inadvertent usage of an adhesive spray used to make greeting cards resulted in vague neurological symptoms and abnormal liver function tests both of which fully resolved on discontinuation.

Keywords: liver function tests; volatile hydrocarbons; inhalation; neurological symptoms.

Elevated transaminases may occur following ingestion and vapour inhalation of some hydrocarbons. We report a case in which inadvertent inhalation resulted in hepatic injury and review the evidence for a link between the hydrocarbons contained in the spray and the patient’s abnormal liver function tests.

Case report
A 63-year-old man attended the Accident and Emergency Department complaining of a 2-month history of non-specific malaise and paraesthesia in his left upper limb. Two days prior to his attendance a general biochemical screen and full blood count were performed by a commercial laboratory through a friend. The results showed an elevated serum alkaline phosphatase of 162 IU/l (normal range 50–136); a serum aspartate transaminase of 73 IU/l (15–37); a serum alanine transaminase of 122 IU/l (30–65) and a serum gamma-glutamyltransferase of 108 IU/l (5–85). The serum bilirubin, albumin and protein were all normal. The patient had been previously well and was not taking any medication. He was a non-smoker and drank 1–2 units of alcohol weekly. He had never previously drunk alcohol excessively. On examination he appeared well, was not jaundiced or pyrexial and had none of the stigmata of chronic liver disease. No hepatomegaly or splenomegaly were detected and the rest of the general examination was unremarkable. Various investigations were arranged and he was discharged from hospital and followed up 3 weeks later in the out-patient clinic. His serological markers for acute viral hepatitis were negative. His serum ceruloplasmin and ferritin level and thyroid function tests were normal. His auto-antibody screen was negative. His abdominal ultrasound revealed an echobright pattern suggestive of mild fatty change and an incidental 2 cm gall stone without biliary tract dilatation. A plain cervical spine radiograph showed minor degenerative changes only.

A week later the patient’s wife, who worked for Age Concern, attended a drug abuse lecture and recognised the potential cause of her husband’s symptoms. He was a keen producer of home-made greeting cards and spent up to 3 hours at a time, two to three times per week, in a poorly ventilated garage using 3M (TM) Spraymount Adhesive. He had been using it regularly for approximately 2 years but because of both a Christmas ‘rush’ to produce extra cards for a charity and excessive cold weather, this use had increased and with virtually no ventilation. The patient then confessed to symptoms of drowsiness and nausea. After stopping it completely and changing to a paste form of adhesive he felt dramatically better with complete cessation of his symptoms. Repeat liver function tests taken 4 weeks after the first set had returned to normal.

References
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