

CONDITIONS CAUSING MALABSORPTION

Coeliac disease

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Coeliac disease is the most common cause of malabsorption in the UK with a reported prevalence of 1:1800, although this may be an underestimate. It is characterised by a hyper-trophic small bowel mucosa with atrophic villi and deep crypts. Loss of surface area and brush border enzymes results in malabsorption. Coeliac disease is caused by an abnormal immune response to gluten, a cereal protein, although the exact mechanism remains unclear. There is a genetic component, as the disease is more common in first-degree relatives and has an association with HLA-B8. In children, coeliac disease presents with steatorrhoea and growth retardation. In adults, it may result in diarrhoea and weight loss but many patients simply present with iron deficiency anaemia. Some patients develop a characteristic skin rash (dermatitis herpetiformis). The diagnosis is usually made after an endoscopic duodenal biopsy allows pathological examination of the mucosa. A blood test for immunoglobulin A anti-tissue transglutaminase (IgA tTGA) is relatively sensitive and specific for diagnosing coeliac disease, making it the preferred test for detection. Measurement of IgA anti-endomysial antibodies (anti-EMA) should be used as a confirmatory test. A duodenal biopsy is usually indicated to confirm the diagnosis. The biopsy usually shows flattening of the mucosa, marked inflammatory changes and characteristic findings of intraepithelial lymphocytes. All tests should be performed while the patient is on a gluten-containing diet as false-negative tests may occur if on a gluten-free diet. Patients with coeliac disease may develop an acute inflammatory condition of the small intestine (ulcerative jejunoileitis) and have an increased risk of small bowel lymphoma and adenocarcinoma. The main treatment for coeliac disease is the withdrawal of gluten from the diet by avoiding wheat, rye and barley. Surgery does not usually play a role in the management of disease and is primarily reserved for resection of malignancy.

(d) (e) Figure 74.9 Common types of blind loop: (a) self-feeding; (b) ileocecal valve incompetence; (c) ileocecal valve stenosis; (d) jejunal diverticula; (e) intestinal stricture causing stasis;

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