

Gluten-Sensitive Enteropathy (Celiac Disease): More Common Than You Think

DAVID A. NELSEN, JR., M.D., M.S., University of Arkansas for Medical Sciences, Little Rock, Arkansas

Gluten-sensitive enteropathy or, as it is more commonly called, celiac disease, is an autoimmune inflammatory disease of the small intestine that is precipitated by the ingestion of gluten, a component of wheat protein, in genetically susceptible persons. Exclusion of dietary gluten results in healing of the mucosa, resolution of the malabsorptive state, and reversal of most, if not all, effects of celiac disease. Recent studies in the United States suggest that the prevalence of celiac disease is approximately one case per 250 persons. Gluten-sensitive enteropathy commonly manifests as "silent" celiac disease (i.e., minimal or no symptoms). Serologic tests for antibodies against endomysium, transglutaminase, and gliadin identify most patients with the disease. Serologic testing should be considered in patients who are at increased genetic risk for gluten-sensitive enteropathy (i.e., family history of celiac disease or personal history of type 1 diabetes) and in patients who have chronic diarrhea, unexplained anemia, chronic fatigue, or unexplained weight loss. Early diagnosis and management are important to forestall serious consequences of malabsorption, such as osteoporosis and anemia. (*Am Fam Physician* 2002;66:2259-66,2269-70. Copyright© 2002 American Academy of Family Physicians.)

C A patient information handout on celiac disease, written by the author of this article, is provided on page 2269.

Although celiac disease was formally described late in the 19th century, treatment remained empiric until the middle of the 20th century when patients were noted to improve dramatically after wheat was removed from their diet. With the development of small-bowel biopsy techniques, the small intestine was identified as the target organ. Disease causality was established when the characteristic features of villous flat-

tening, crypt hyperplasia, and increased intraepithelial lymphocytes (*Figure 1*) were shown to normalize after the institution of a gluten-free diet.¹

In the mid-1960s, an enteropathy strikingly similar to celiac disease was identified in patients with dermatitis herpetiformis. Subsequently, this skin disorder was shown to be a manifestation of gluten-sensitive enteropathy. In the mid-1960s, adult celiac disease was also noted to be associated with numerous neuro-

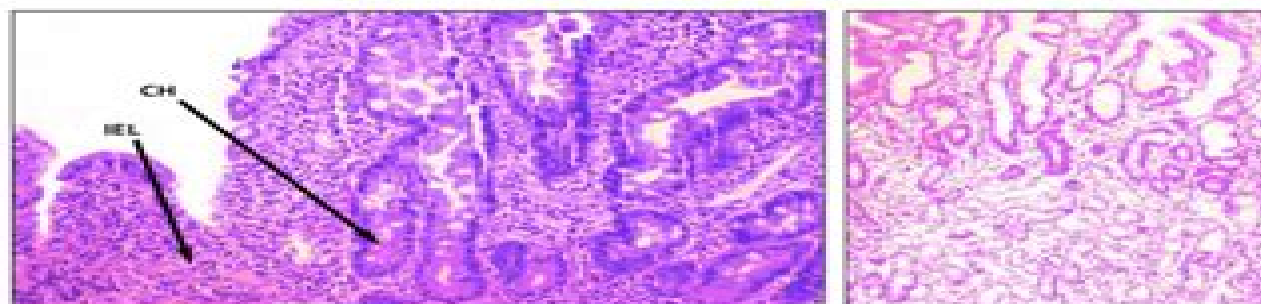


FIGURE 1. (Left) Photomicrograph of distal duodenal biopsy specimen in a patient with celiac disease. Note the characteristic features of crypt hyperplasia (CH), and increased intraepithelial lymphocytes (IEL). (Right) For comparison, a normal biopsy specimen is shown.

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