

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

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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 I 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.)

 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-

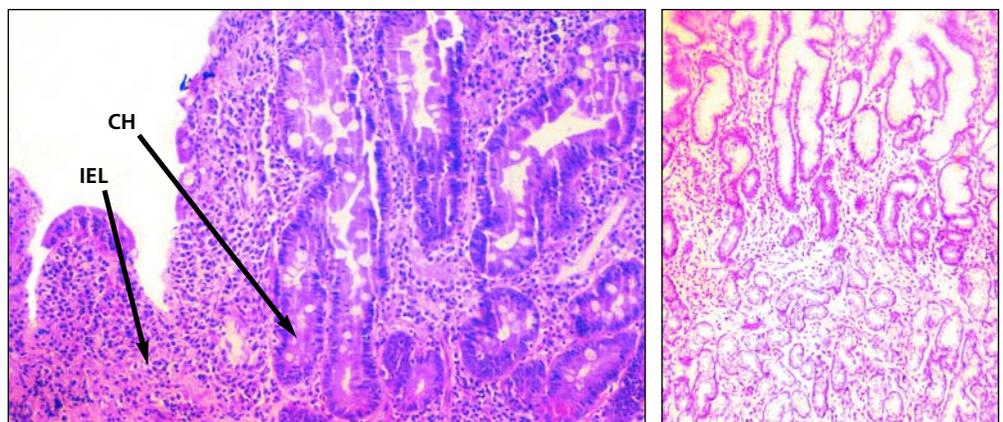


FIGURE 1. (Left) Photomicrograph of distal duodenal biopsy specimen in a patient with celiac disease. Note the characteristic features of crypt hyperplasia (CH), and increased intraepithelial lymphocytes (IEL). (Right) For comparison, a normal biopsy specimen is shown.

logic disorders, including epilepsy, cerebral calcifications, and peripheral neuropathy.¹

Recent population studies indicate that celiac disease is more common than was previously thought. Some patients with gluten-sensitive enteropathy have minimal or no symptoms and are unlikely to be referred to a gastroenterologist unless the disease is considered. Hence, family physicians need to be familiar with the diagnosis and management of gluten-sensitive enteropathy.

Pathophysiology

Ingested protein does not normally provoke an immune response. This phenomenon is termed "oral tolerance." Patients who exhibit true allergy to an ingested protein (e.g., milk or soy protein) have a typical IgE-mediated response consisting of urticaria, angioedema, and bronchoreactivity.

The autoimmunity in gluten-sensitive enteropathy involves plasma cells that produce IgA and IgG; there is little or no IgE involvement. Current theory suggests that ingested α -gliadin (a component of the gluten protein) and related peptides bind with tissue transglutaminase (a ubiquitous intracellular enzyme) in enterocytes. The α -gliadin is rich in glutamine; transglutaminase deamidates glutamine residues, forming glutamic acid. Deamidation enhances the immunogenicity of α -gliadin by creating epitopes that are recognized as foreign by host cell-mediated immunity.²

Plasma cells produce IgA and IgG that are directed against a variety of antigens, including transglutaminase, endomysium, gliadin, and reticulon. Locally elaborated lymphokines attract inflammatory cells.³ This intense

TABLE 1
Celiac Disease and Other Autoimmune Disorders*

Type 1 diabetes mellitus	Recurrent aphthous ulcerations
Autoimmune thyroid disease	Sjögren's syndrome
Rheumatoid arthritis	Sarcoidosis
Systemic lupus erythematosus	Vitiligo or alopecia areata
Autoimmune hepatitis	IgA deficiency
Autoimmune Addison's disease	Psoriasis†

*—Listed in descending order of clinical importance and prevalence.

†—Observed association.

local inflammatory reaction produces the villous flattening characteristic of gluten-sensitive enteropathy. Malabsorption of micronutrients (e.g., vitamins and minerals) and macronutrients (e.g., protein, carbohydrate, fat) follows. Small-bowel involvement is most prominent proximally and may be "patchy," especially in patients with "silent" celiac disease (i.e., minimal or no symptoms) and those with dermatitis herpetiformis.

Approximately 95 percent of patients with celiac disease exhibit specific Human Leukocyte Antigen (HLA) class II alleles DQA1*0501 and DQB1*0201.⁴ Patients with type 1 diabetes, autoimmune thyroid disease,⁵ Sjögren's syndrome, primary biliary cirrhosis, Addison's disease, systemic lupus erythematosus, selective IgA deficiency, and alopecia areata may also exhibit similar genotypes and are at risk for gluten-sensitive enteropathy (Table 1). Because many persons have these genotypes and only a few develop gluten-sensitive enteropathy, investigators have hypothesized that other genes or cofactors may be involved.¹

Epidemiology

Until recently, celiac disease was considered to be relatively uncommon. Previous U.S. figures suggested that it affected one in 6,000 persons.⁶ However, population studies published in the past four years suggest a much higher prevalence, particularly in persons of European ancestry.⁷ Studies conducted in Europe estimate the seroprevalence of celiac disease to be one case per 130 to 300 persons.⁸⁻¹⁰

In a recent U.S. study,¹¹ investigators tested sera from 2,000 healthy Red Cross blood donors and found eight samples that were positive for antibodies associated with gluten-sensitive enteropathy (seven samples from white

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TABLE 2

Symptoms of Celiac Disease and Possible Causes

<i>Symptoms</i>	<i>Possible causes</i>
Fatigue, malaise	Anemia, general immune system activation
Weight loss	Nutrient malabsorption
Diarrhea, abdominal pain	Accelerated gastrointestinal tract transit time, steatorrhea, malabsorption
Anemia	Most commonly, iron deficiency; less commonly, vitamin B ₁₂ and/or folate deficiency
Bone pain	Osteoporosis
Aphthous oral ulcers, glossitis, stomatitis	Vitamin deficiency, "oral" celiac disease
Infertility	Postulated cause: iron, folate, and/or zinc deficiency
Male impotence, decreased libido	Peripheral insensitivity to circulating testosterone
Alopecia areata	Immunologic attack on hair follicles
Dental enamel defects	Deminceralization during tooth bud development in children
Hypoglycemia	Delayed absorption of glucose
Gas, flatus, borborygmus	Secondary digestion of sugars by intestinal flora
Seizures, gluten ataxia, central nervous system symptoms	Increased affinity of celiac antibodies for brain vasculature

persons, one sample from a black person). The seroprevalence rate in this study (one case per 250 persons tested) is consistent with the rates in European studies.

The likelihood of having gluten-sensitive enteropathy increases to 10 to 20 percent in persons who have a first-degree relative with celiac disease.¹ In addition, celiac disease is associated with other autoimmune syndromes. For example, as many as 7 percent of patients with type I diabetes also have gluten-sensitive enteropathy.¹²

Clinical Presentation

Untreated gluten-sensitive enteropathy is associated with a range of symptoms^{13,14} (Table 2). The "classic" form typically presents in infancy and manifests as failure to thrive, diarrhea, abdominal distention, developmental delay, and, occasionally, severe malnutrition. Failure to diagnose the disorder may lead to a true medical emergency.

Beyond infancy, the symptoms of celiac disease tend to be less dramatic. Older children may present with constitutional short stature or dental enamel defects.

Women comprise approximately 75 percent of newly diagnosed adult celiac disease cases. Women also tend to have more clinically conspicuous disease.¹⁵ In adults of both sexes, gastrointestinal tract involvement may mani-



FIGURE 2. Dermatitis herpetiformis, sometimes termed "celiac disease of the skin." Vesicular, crusted, intensely pruritic lesions develop on the back (*top left and right*), buttocks (*lower left*), and elbows (*lower right*). Secondary infection of the lesions is common.

fest as diarrhea, constipation, or other symptoms of malabsorption, such as bloating, flatus, or belching. Fatigue, depression, fibromyalgia-like symptoms, aphthous stomatitis, bone pain, dyspepsia, gastroesophageal reflux, and other nonspecific symptoms may be present and can make the diagnosis quite challenging.¹⁴ A number of other autoimmune syndromes have been associated with celiac disease (Table 1).

DERMATITIS HERPETIFORMIS

Fewer than 10 percent of adults with gluten-sensitive enteropathy present with dermatitis herpetiformis.¹⁶ This skin condition may be misdiagnosed as atypical psoriasis or nonspecific dermatitis.

The rash of dermatitis herpetiformis is intensely pruritic and typically occurs on the back, buttocks, knees, and elbows (Figure 2). Unexcoriated lesions (Figure 3) are

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FIGURE 3. Close-up view of dermatitis herpetiformis. Uncoriated lesions are remarkably like those of herpes simplex (thus, the term “herpetiformis”).

remarkably like those of herpes simplex (thus, the term “herpetiformis”). Granular IgA deposition on immunofluorescence of a skin biopsy specimen is diagnostic (Figure 4).

ANEMIA

Anemia is the most common laboratory manifestation of celiac disease^{13,14} (Table 3). One half of patients with newly diagnosed gluten-sensitive enteropathy are anemic. Iron is absorbed in the proximal small intestine, where celiac manifestations are most prominent; hence, iron malabsorption is common. In addition, occult blood loss related to intense small-bowel inflammation may occur in 50 percent of patients with gluten-sensitive enteropathy.¹⁷ Less commonly, vitamin B₁₂ deficiency, folate deficiency, or both may be present.

SILENT CELIAC DISEASE

A number of investigators believe that clinically apparent gluten-sensitive enteropathy represents the “tip of the iceberg” of the overall disease burden.⁷ Patients who were detected in the seroprevalence studies⁸⁻¹¹ were asymptomatic or oligosymptomatic (so-called “silent” celiac disease).

Family physicians should consider serologic testing in patients with the following: family history of celiac dis-

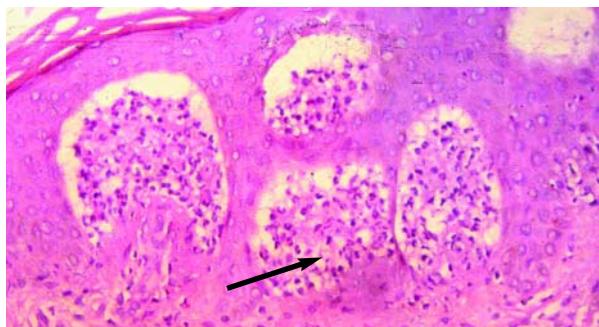


FIGURE 4. Photomicrograph of a skin biopsy specimen in a patient with dermatitis herpetiformis. Note the intense inflammatory infiltration at the dermoepidermal junction. Immunofluorescence demonstrated granular deposition of IgA (arrow).

ease, personal history of thyroid disease or type I diabetes, irritable bowel syndrome, anemia (especially iron deficiency), chronic diarrhea, chronic fatigue, unexplained weight loss, short stature, epilepsy, infertility,¹⁸ or unexplained elevation of transaminase levels. Asymptomatic or oligosymptomatic patients are still at risk for complications of celiac disease.

Diagnosis

SEROLOGIC TESTS

When the diagnosis of gluten-sensitive enteropathy is suspected, serologic tests can identify many affected patients.¹⁹ (Figure 5). IgA antiendomysial antibody has been shown to be 85 to 100 percent sensitive and 96 to 100 percent specific for celiac disease. IgA antiendomysial antibody is measured using direct immunofluorescence of monkey esophagus or umbilical cord tissue processed with suspect serum.

IgG and IgA antigliadin antibodies are also useful in diagnosing celiac disease. The antibody directed against endomysium has recently been identified as identical to the antibody directed against (tissue) transglutaminase. An enzyme-linked immunosorbent assay has been developed to measure IgA antitransglutaminase; this test will generally replace the more tedious direct fluorescent antibody test for IgA antiendomysial antibody.²⁰⁻²² In diagnosing celiac disease, antitransglutaminase antibody is considered to be approximately as sensitive and specific as antiendomysial antibody (Table 4).^{21,23-25}

The presence of IgA antiendomysial and antitransglutaminase antibodies correlates with intestinal damage. Tests for these antibodies are highly sensitive in patients with total

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TABLE 4
Serologic Tests for Celiac Disease

<i>Antibody test</i>	<i>Sensitivity (%)</i>	<i>Specificity (%)</i>	<i>Time course</i>	<i>Cost*</i>
IgA antiendomysial antibody	85 to 100	96 to 100	Antibody disappears within several months after institution of gluten-free diet.	\$45 to 99
IgA antitransglutaminase antibody	95	90	Limited data; correlated with IgA antiendomysial antibody in studies	85 to 164
IgA anti gliadin antibody	53 to 100	65 to 100	More persistent than IgA antiendomysial antibody; may persist for 6 months or longer	45
IgG anti gliadin antibody	57 to 100	42 to 98	Most persistent; may be detectable up to 12 months after institution of gluten-free diet False-positive tests reported in patients with Crohn's disease, wheat-protein allergy, and postdiarrhea states	45 to 90

*—Average of costs at three reference laboratories in the summer of 2001.

Information from references 21 through 23 and 25

Declining autoantibody titers correlate with resolution of the gastrointestinal lesion and may be used to document clinical improvement. In equivocal cases or when a patient has been on a gluten-free diet, a gluten challenge test may be used to provoke the gastrointestinal lesion and serologic response.

DISTAL DUODENAL BIOPSY

Distal duodenal biopsy is the gold standard for the diagnosis of celiac disease. Biopsy should be performed in most patients with suspected gluten-sensitive enteropathy. In determining the need for biopsy, family physicians should consult a gastroenterologist who is experienced in the diagnosis and management of celiac disease.¹

Complications

OSTEOPOROSIS

Calcium and vitamin D malabsorption dramatically increases the risk of osteoporosis and osteomalacia in patients with gluten-sensitive enteropathy. Most patients with celiac disease have some degree of osteopenia or osteoporosis. Fortunately, calcium and vitamin D supplementation, coupled with a strict gluten-free diet, usually results in remineralization of the skeleton.^{27,28}

Bone density should be assessed in all patients with newly diagnosed celiac disease. Postmenopausal women should be advised about the benefits of estrogen replacement. The use of antiresorptive agents (e.g., alendronate [Fosamax]) has not been extensively studied, but these drugs may be of benefit.

NEUROLOGIC MANIFESTATIONS

Cerebral calcifications and epilepsy have been associated with celiac disease^{29,30} and do not always resolve with

the institution of a gluten-free diet. Peripheral neuropathy, postural instability, "gluten ataxia," and other vague neurologic complaints may be the sole manifestation of gluten-sensitive enteropathy.³¹ Autoantibodies associated with celiac disease have demonstrated a strong affinity for brain vasculature.³²

REFRACTORY SPRUE

In patients with refractory sprue, gastrointestinal tract inflammation continues despite maintenance of a gluten-free diet. Dietary noncompliance is the most common reason for persistent inflammation; however, coexistent conditions such as hyperthyroidism and collagenous colitis should also be considered.

Patients who are truly refractory to dietary measures may have cryptic lymphoma of (bowel) intraepithelial lymphocytes. All diet-refractory patients should be evaluated by a gastroenterologist with expertise in the management of celiac disease. Corticosteroids and immunosuppressant drugs have been used to treat refractory sprue, but data on their effectiveness are sparse.

Intestinal strictures and bowel obstruction may develop in patients with refractory sprue or celiac disease that has been untreated over a long period.

LYMPHOMA AND BOWEL ADENOCARCINOMA

Enteropathy-associated T-cell lymphoma has been associated with untreated gluten-sensitive enteropathy and refractory sprue.³³ Lymphoma may develop in patients with celiac disease who also have dermatitis herpetiformis.

Studies^{34,35} have shown that maintenance of a long-term gluten-free state reduces the risk of lymphoma to the level in the general population. Thus, it is imperative that

TABLE 5

Laboratory Evaluation of Patients with Newly Diagnosed Celiac Disease

Hematology

Complete blood cell count
Platelet count

Laboratory tests

Iron level, total iron-binding capacity determination, ferritin level*
Vitamin B₁₂ and folate levels
Calcium and phosphate levels
Alkaline phosphatase level
Blood urea nitrogen and creatinine levels
Albumin and total serum protein levels
Aspartate transaminase and alanine transaminase levels

Imaging

Dual energy x-ray absorptiometry (DEXA) of spine and hip

Serologic tests†

Quantitative IgA antiendomysial antibody or quantitative IgA antitransglutaminase
Quantitative IgA and IgG antigliadin antibodies

*—The ferritin level may be misleading: it may be elevated as an acute-phase reactant.

†—Serologic testing should be recommended for first-degree relatives of patients with newly diagnosed celiac disease.

patients with celiac disease (and dermatitis herpetiformis) maintain a gluten-free diet for the rest of their lives.

Patients with celiac disease are also at risk for the development of bowel adenocarcinoma in all sites. The risk is especially high in patients with a long period of disease preceding institution of a gluten-free diet.

Management

Once the diagnosis of celiac disease has been made, patients should be evaluated for known manifestations and complications (Table 5). Iron deficiency should be treated with supplemental iron. Osteoporosis should be treated with calcium and vitamin D replacement. Depending on individual factors, patients with gluten-sensitive enteropathy may need to take a multivitamin, iron, calcium, magnesium, zinc, selenium, vitamin D, or other nutrients.

The primary treatment for celiac disease is the removal of gluten and related proteins from the diet. Complete exclusion of dietary gluten generally results in rapid and complete healing of small-bowel inflammation. Advice from a registered dietitian is essential to outline an appropriate diet.

Gluten, a prolamine, is the primary protein in wheat. Hence, wheat and products containing wheat must be avoided. Barley and rye contain similar proteins and must also be avoided. Oats are a subject of controversy.^{36,37}

Although oats themselves may be nontoxic in limited quantities, commercial oat products are measurably contaminated with wheat. Rice, corn, maize, flax, quinoa, tapioca, potato, amaranth, and other grain substitutes, such as nuts and beans, are safe.

There are many commercial gluten-free products, including breads, cookies, chips, and cereals, that can be used to fashion a rich and interesting diet. Meats, vegetables, fruit, and most dairy products are free of gluten, as long as they have not been contaminated during production.

A number of food manufacturers maintain lists of gluten-free products. These lists can be obtained from the manufacturers' Web sites or by telephone request. Information on gluten-free foods is also available from local or national support groups such as the Celiac Sprue Association. Selected resources are provided in Table 6.

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Figure 1 provided by Laura Lamps, M.D., assistant professor of pathology, University of Arkansas for Medical Sciences (UAMS), Little Rock. Figure 4 provided by Bruce Smoller, M.D., professor of pathology and dermatology, UAMS.

TABLE 6

Selected Resources

Celiac Sprue Association/United States of America, Inc.*

P.O. Box 31700, Omaha, NE 68131-0700
Telephone: 402-558-0600
Fax: 402-558-1347
Web site: www.csaceliacs.org
E-mail address: celiacs@csaceliacs.org

Celiac Discussion List Archives

Web site: www.fastlane.net/homepages/thodge/archive.shtml

National Digestive Diseases Information Clearinghouse

Web site: www.niddk.nih.gov/health/digest/pubs/ceciac/index.htm

Celiac Disease Resources for Medical Professionals†

Web site: www.uams.edu/ceciac

*—The Celiac Sprue Association is a national organization that supports persons with celiac disease and related conditions.

†—David A. Nelsen, Jr., M.D., M.S., maintains a Web site with resources for primary care physicians to assist them in the care of patients with celiac disease.

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