

A young woman with fatigue

22-YEAR-OLD WOMAN presents to the Clinic for evaluation of fatigue. She has not felt well for the past few years. Her current symptoms include generalized fatigue and diarrhea, characterized as two to three semiformed, nonbloody bowel movements each day and occasional episodes of watery diarrhea. Her bowel movements are usually precipitated by meals. She consumes a regular diet and has not recognized any intolerance to any particular foods. She denies having any abdominal pain, nausea, vomiting, recent travel, joint pain, rash, or change in the texture of her hair. She has been seen by several internists in her hometown, who have not provided her with a specific diagnosis.

Her medical history is significant for anemia, anxiety, and depression. Menarche occurred at age 16. Her menstrual cycle has been regular, with bleeding noted to be only modest. Her medications include oral contraceptive pills. She has not had previous surgeries.

On examination, she appears well. She is afebrile, weighs 128 lbs, and is 63 inches tall. The physical examination is normal, including a rectal examination and fecal occult blood testing.

Routine laboratory tests are performed. Results:

- White blood cell count $3.88 \times 10^9/L$ (normal 4.0–11)
- Hemoglobin 10.4 g/dL (normal 12–16)
- Hematocrit 34% (normal 37%–47%)
- Mean corpuscular volume 80.2 fL (normal 80–100)
- Mean corpuscular hemoglobin 24.5 pG (normal 27–34)
- Platelet count 365 x 109/L (normal 150–400)
- Sodium 141 mmol/L (normal 132–148)
- Potassium 4.2 mmol/L (normal 3.5–5.0)

- Chloride 107 mmol/L (normal 98–110)
- Alanine aminotransferase 22 U/L (normal 0–45)
- Glucose 66 mg/dL (normal 65–100)
- Blood urea nitrogen 6 mg/dL (normal 8–25)
- Creatinine 0.6 mg/dL (normal 0.7–1.4)
- Thyroid-stimulating hormone 2.860 mIU/L (normal 0.4–5.5)
- Red blood cell folate 539 ng/mL (normal 257–800)
- Vitamin B₁₂ 321 pg/mL (normal 221–700)
- Iron/total iron-binding capacity 21/445 µg/dL (normal 30–140, 210–415)
- Ferritin 5 ng/mL (normal 9–150).

DIFFERENTIAL DIAGNOSIS

- 1 Which of the following is the most likely cause of her diarrhea?
- ☐ Thyroid disease
- ☐ Functional bowel disease
- ☐ Gluten-sensitive enteropathy (celiac disease)

Given her constellation of symptoms (fatigue, neuropsychiatric changes, iron deficiency anemia, and diarrhea), celiac disease is the most likely diagnosis. Hyperthyroidism can cause diarrhea, but this is unlikely since her thyroid tests are normal. Functional bowel disease is a diagnosis of exclusion and usually has a more chronic, fluctuating course.

CELIAC DISEASE HAS VARIOUS PRESENTATIONS

Celiac disease has various presentations and therefore has been classified into several types^{1,2}:

Classic disease is dominated by symptoms of malabsorption. The diagnosis is established

Atypical presentations of celiac disease are now more common than classic disease in adults

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by serologic testing, findings of villous atrophy on biopsy, and improvement of symptoms on a gluten-free diet. However, the presentation of celiac disease has changed, and now atypical presentations are more common in adults (see below). The reason for the change in presentation is not known, but some have hypothesized that it is related to an increase in breastfeeding and the later introduction of cereals into infants' diets.

Celiac disease with atypical symptoms is characterized by extraintestinal manifestations with few or no gastrointestinal (GI) symptoms. Patients may present with irondeficiency anemia; osteoporosis or vitamin D deficiency; arthritis; neurologic symptoms such as ataxia, headaches, or depression or anxiety; myocarditis; infertility; or elevated aminotransferase levels. As in classic celiac disease, the diagnosis is established with serologic testing, findings of villous atrophy on biopsy, and improvement of symptoms on a gluten-free diet.

Latent disease includes cases in patients with positive serologic tests but no villous atrophy on biopsy. These patients have no symptoms but may develop symptoms or histologic changes later.

Silent disease refers to cases in patients who have no symptoms but have a positive serologic test and villous atrophy on biopsy. These cases are usually detected via screening of people at high risk, ie, relatives of patients with celiac disease.

It is important that clinicians be aware of the various symptoms and presentations of celiac disease in order to make the diagnosis.

CONFIRMING CELIAC DISEASE

- **2** Which of the following is used to test for celiac disease?
- ☐ Immunoglobulin G (IgG) and immunoglobulin A (IgA) antigliadin antibody testing
- ☐ IgA antiendomysial antibody and IgA antitransglutaminase antibody testing
- ☐ HLA DQ2/DQ8 testing

The sensitivity of antigliadin antibody testing is only about 70% to 85%, and its specificity is about 70% to 90%. Better serologic tests are

those for IgA antiendomysial and antitransglutaminase antibodies, which have sensitivities greater than 90% and specificities greater than 95%.³ HLA DQ2/DQ8 testing has a high sensitivity (> 90%–95%), but because about 30% of the general population also carry these markers, the specificity of this test is not ideal. This test is best used for its negative predictive value—ie, to rule out the diagnosis of celiac disease.

Of note: 1% to 2% of patients with celiac disease have a deficiency of IgA.⁴ Therefore, if the clinical suspicion for celiac disease is high but the IgA antibody tests are negative or equivocal, IgG antitransglutaminase and IgG antiendomysial antibody tests can help establish the diagnosis. HLA testing in this situation can also help rule out the diagnosis.

CONFIRMING CELIAC DISEASE— CONTINUED

- **3** What test should be performed next in this patient?
- ☐ Upper GI series with small-bowel follow-through
- Esophagogastroduodenoscopy with biopsies
- ☐ Small-bowel barium study
- ☐ Video capsule endoscopy

Today, the presumptive diagnosis of celiac disease requires positive serologic testing and biopsy results. Esophagogastroduodenoscopy with biopsies should be ordered. Upper GI series and barium studies do not provide a tissue diagnosis. Barium studies and other radiologic tests can be considered if a patient does not have the expected response to a strict gluten-free diet or if one suspects complications of celiac disease, such as GI lymphoma.

Video capsule endoscopy is an emerging tool for diagnosing celiac disease, as suggested in several trials.⁵ Some findings seen on video capsule endoscopy in patients with celiac disease include mosaicism, nodularity, visible vessels, and loss of mucosal folds. However, the role of this test continues to be investigated, and biopsy is still required to confirm the diagnosis.

A definitive diagnosis is confirmed when symptoms resolve with a gluten-free diet, and repeat biopsies (3–9 months after dietary changes) show histologic improvement,

All patients with celiac disease should receive dietary counseling

Small-bowel biopsy findings: Normal

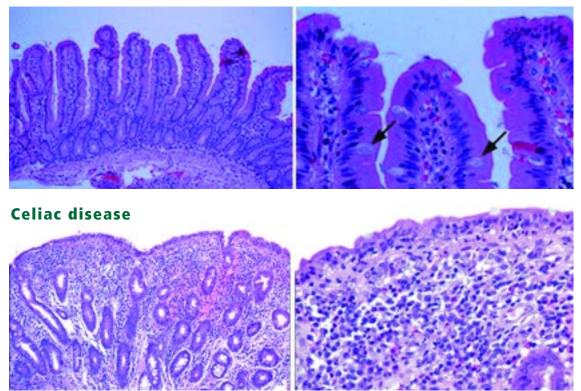


FIGURE 1. A low-power view (top left) and high-power view (top right) of the normal villous architecture of the small intestine. The high-power view shows the enterocytes and interspersed goblet cells (arrows). Bottom left, a sample from a patient with celiac disease shows severe villous blunting in this low-power view. Bottom right, a higher-power view shows loss of columnar epithelial shape, mucin depletion, and an increased nuclear-tocytoplasmic ratio within the epithelial layer. Within the normal epithelial layer, there is an increased ratio of lymphocytes to epithelial cells. The normal ratio of lymphocytes to epithelial cells is 1 to 5-10. In this specimen the ratio is approximately 3 to 5 lymphocytes to 5 epithelial cells. The lamina propria also shows an increased number of plasma cells. This constellation of microscopic findings is characteristic of gluten-sensitive enteropathy.

Celiac disease may have a patchy distribution. so multiple biopsy samples should be taken

although the need for repeat biopsy is controversial. A hallmark histologic finding is villous atrophy (FIGURE 1). However, villous atrophy may be patchy, and it is recommended that multiple biopsy specimens be taken from the duodenal mucosa to increase the diagnostic yield.

WHO SHOULD BE TESTED FOR CELIAC DISEASE?

The reported prevalence of symptomatic celiac disease is about 1 in 1,000 live births in populations of northern European ancestry, ranging from 1 in 250 (in Sweden) to 1 in

4,000 (in Denmark).6 The prevalence appears to be higher in women than in men.⁷

In a large US study, the prevalence of celiac disease was 1 in 22 in first-degree relatives of celiac patients, 1 in 39 in second-degree relatives, 1 in 56 in patients with either GI symptoms or a condition associated with celiac disease, and 1 in 133 in groups not at risk.8 Another study found that the prevalence of antiendomysial antibodies in US blood donors was as high as 1 in 2,502.

Given that patients with celiac disease may not present with classic symptoms, it has been suggested that the following groups of patients be tested for it1:

- Patients with GI symptoms such as chronic diarrhea, malabsorption, weight loss, or abdominal symptoms
- Patients without diarrhea but with other unexplained signs or symptoms that could be due to celiac disease, such as iron-deficiency anemia, elevated aminotransferase levels, short stature, delayed puberty, or infertility
- Symptomatic patients at high risk for celiac disease. Risk factors include type 1 diabetes or other autoimmune endocrinopathies, first- and second-degree relatives of people with celiac disease, and patients with Turner, Down, or Williams syndromes.

Screening of the general population is not recommended, even in populations at high risk (eg, white people of northern European ancestry).

■ WHAT CAN CELIAC PATIENTS EAT?

- **4** Patients with celiac disease should avoid eating which of the following?
- Wheat
- □ Barley
- □ Rye
- ☐ Oats

Patients with celiac disease should follow a gluten-free diet and should initially eliminate all of these substances.

Some recent studies have suggested that pure oat powder can be tolerated without disease recurrence, although the long-term safety of oat consumption in patients with celiac disease is uncertain. It may be reasonable for patients to reintroduce oats when the disease is under control, especially since uncontaminated oats can be obtained from reliable retail or wholesale stores. The definitive diagnosis of celiac disease requires clinical suspicion, serologic tests, biopsy, and documented clinical and histologic improvement after a gluten-free diet is started.

All patients with celiac disease should receive dietary counseling and referral to a nutritionist who is experienced in the treatment of this disease. Because of the significant lifestyle and dietary changes involved in treating this disease, many patients may also benefit from participating in a celiac support group.

COMPLICATIONS OF CELIAC DISEASE

- ☐ Anemia
- Osteoporosis
- ☐ Intestinal lymphoma
- ☐ Infertility
- Neuropsychiatric symptoms
- □ Rash

All of the above are complications of untreated celiac disease and are often clinical features at presentation. Patients with celiac disease should be tested for anemia and nutritional deficiencies, including iron, folate, calcium, and vitamin D deficiency.

All patients should also undergo dualenergy x-ray absorptiometric scanning. Bone loss is thought to be related to vitamin D deficiency and secondary hyperparathyroidism, and may be partially reversed with a glutenfree diet.

Celiac disease is associated with hyposplenism, so pneumococcal vaccination should be considered. Celiac disease is also frequently associated with the rash of dermatitis herpetiformis, and diagnosis of this rash should prompt an evaluation for celiac disease.

Other associated conditions include Down syndrome, selective IgA deficiency, and other autoimmune diseases such as type 1 diabetes, thyroid disease, and liver disease.

■ WHAT HAPPENED TO OUR PATIENT?

Our patient tested positive for antiendomysial and antitransglutaminase antibodies and underwent small-bowel biopsy, which confirmed the diagnosis of celiac disease. She was started on a gluten-free diet, and within 2 weeks she noted an improvement in her symptoms of fatigue, GI upset, mood disorders, and difficulty with concentration. She met with a nutritionist who specializes in celiac disease and joined a celiac support group.

However, about 2 months later, her symptoms recurred. She again met with her nutritionist, who confirmed that she was adhering to a gluten-free and lactose-free diet. Even so, when she was tested again for

Screening for celiac disease in the general population is not recommended

antitransglu-taminase antibodies, the titer was elevated. Stool cultures were obtained and were negative. She was started on a course of prednisone, and her symptoms resolved.

WHAT IF PATIENTS DO NOT RESPOND TO TREATMENT?

The most common cause of recurrent symptoms or nonresponse to treatment is noncompliance with the gluten-free diet or inadvertent ingestion of gluten. Patients who do not respond to treatment or who have a period of response but then relapse should be referred back to a nutritionist who specializes in celiac disease.

If a patient continues to have symptoms despite strict adherence to a gluten-free diet, other disorders should be considered, such as concomitant lactose intolerance, small-bowel bacterial overgrowth, pancreatic insufficiency, or irritable bowel syndrome. If these conditions are ruled out, patients can be considered for treatment with prednisone or other immunosuppressive agents. Patients with refractory symptoms are at higher risk of more severe complications of celiac disease, such as intestinal lymphoma, intestinal strictures, and collagenous colitis.

TAKE-HOME POINTS

Celiac disease classically presents with symptoms of malabsorption, but nonclassic presentations are much more common.

REFERENCES

- 1. National Institutes of Health. NIH Consensus Development Conference on Celiac Disease, 2004. http:// consensus.nih.gov/2004/2004CeliacDisease118html.htm. Accessed 1/29/2008.
- 2. Rostom A, Murray JA, Kagnoff MF. American Gastroenterological Association (AGA) Institute technical review on the diagnosis and management of celiac disease. Gastroenterology 2006; 131:1981-2002.
- 3. Hellekson K. AHRQ releases practice guidelines for celiac disease screening. Am Fam Phys 2005: 71:1-3.
- 4. Cataldo F, Marino V, Bottaro G, Greco P, Ventura A. Celiac disease and selective immunoglobulin A deficiency. J Pediatr 1997; 131:306-308.
- 5. Kesari A, Bobba RK, Arsura EL. Video capsule endoscopy and celiac disease. Gastrointest Endosc 2005; 62:796-797.
- 6. Branski D, Fasano A, Troncone R. Latest developments in

- Celiac disease should be tested for in patients with or without symptoms of malabsorption and other associated signs or symptoms including unexplained irondeficiency anemia, infertility, short stature, delayed puberty, or elevated transaminases. Testing should be considered for symptomatic patients with type 1 diabetes or other autoimmune endocrinopathies, first- and second-degree relatives of patients with known disease, and those with certain chromosomal abnormalities.
- Heightened physician awareness is important in the diagnosis of celiac disease.
- Diagnosis depends on serologic testing, biopsy, and clinical improvement on a gluten-free diet.
- Treatment should consist of education about the disease, consultation with a nutritionist experienced in celiac disease, and lifelong adherence to a gluten-free diet. Referral to a celiac support group should be considered.
- Long-term follow-up should include heightened vigilance and awareness of the complications of celiac disease such as osteoporosis, vitamin D deficiency and other nutritional deficiencies, increased risk of malignancy, association with low birth-weight infants and preterm labor, and occurrence of autoimmune disorders.

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Test patients for anemia and iron, folate, calcium, and vitamin D deficiency

- the pathogenesis and treatment of celiac disease. J Pediatr 2006; 149:295-300.
- 7. Rampertab SD, Pooran N, Brar P, Singh P, Green PH. Trends in the presentation of celiac disease. Am J Med 2006; 119(4):355.e9-e14.
- 8. Fasano A, Berti I, Gerarduzzi T, et al. Prevalence of celiac disease in at-risk and not-at-risk groups in the United States: a large multicenter study. Arch Intern Med 2003; 163:286-292.
- 9. Janatuinen EK, Pikkarainen PH, Kemppainen TA, et al. A comparison of diets with and without oats in adults with celiac disease. N Engl J Med 1995; 333:1033-1037.

ADDRESS: Kathryn A. Teng, MD, Department of General Internal Medicine, S70, Cleveland Clinic, 9500 Euclid Avenue, Cleveland, OH 44195; e-mail tengk@ccf.org.

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