Case report

Navigating a Complex Clinical Conundrum: The Coexistence of Crohn’s Disease, Celiac Disease, and Ankylosing Spondylitis

.ABSTRACT

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| The coexistence of Crohn’s disease (CrD), celiac disease (CD), and ankylosing spondylitis (AS) is exceptionally rare and poses significant diagnostic and therapeutic challenges. We report the case of a 27-year-old woman with alpha-thalassemia and a nonspecific genetic disorder, who presented with chronic diarrhea, severe weight loss, and Koenig syndrome. Laboratory results confirmed malabsorption syndrome, anemia, hypoalbuminemia, and elevated inflammatory markers, with colonoscopy and biopsies diagnosing CrD and CD. Six months later, she developed peripheral arthritis, leading to a diagnosis of AS. Treatment with corticosteroids, azathioprine, a gluten-free diet, and infliximab led to clinical remission. This case underscores the importance of thorough diagnostic evaluation and highlights the effectiveness of infliximab in managing overlapping autoimmune conditions. |

*Keywords: Crohn-Celiac-Ankylosing Spondylitis*

1. INTRODUCTION

Celiac disease (CD) is a chronic systemic autoimmune disorder that affects genetically predisposed individuals and is triggered by the ingestion of gluten. It is characterized by histologically confirmed villous atrophy, lymphocytic infiltration, and clinical improvement following a gluten-free diet (GFD) [1]. On the other hand, Crohn’s disease (CrD) is a chronic inflammatory disorder of the gastrointestinal tract, which can involve any part of the digestive system, but most frequently affects the terminal ileum and proximal colon. The coexistence of these two diseases is rare, and reports in the literature are limited.

2. Case presentation

We present the case of a 27-year-old woman with a history of alpha-thalassemia and a nonspecific genetic disorder. She was referred for evaluation due to a six-month history of chronic diarrhea (approximately six stools per day), significant weight loss (20 kg), and abdominal pain consistent with Koenig syndrome. On initial physical examination, the patient appeared pale and emaciated, with a weight of 34 kg and a height of 1.56 meters (BMI: 14 kg/m²). No peripheral lymphadenopathy or abdominal masses were noted.

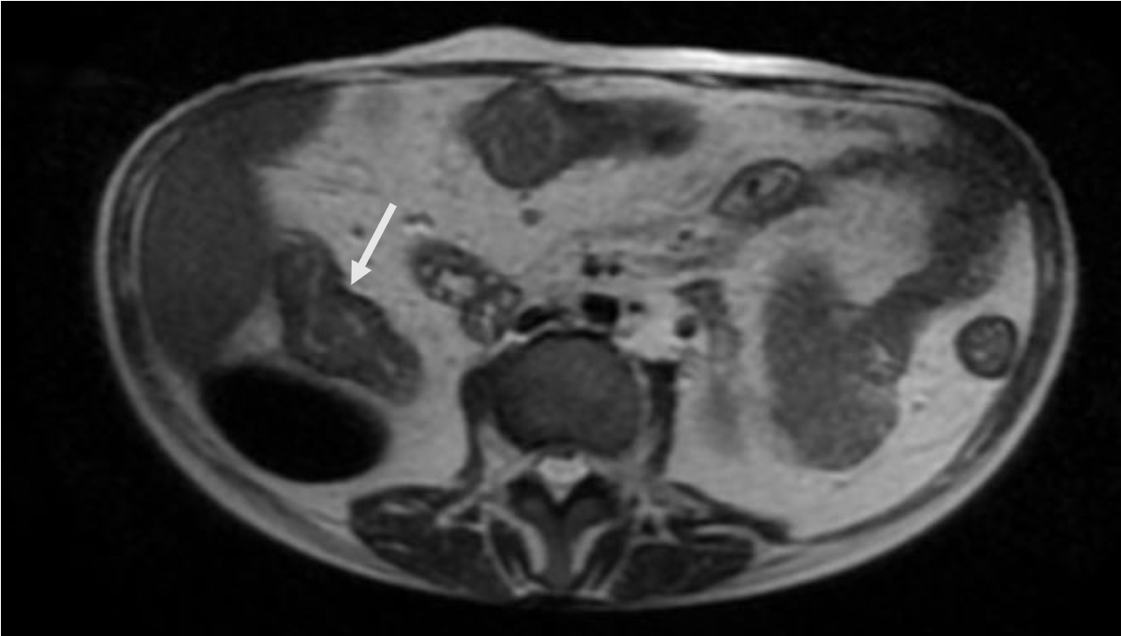
**Medical and Surgical History**The patient had no prior history of gastrointestinal surgery or other chronic illnesses except for alpha-thalassemia. Her family history was unremarkable for autoimmune diseases or gastrointestinal disorders. She denied previous hospitalizations or significant medical interventions, apart from routine management of alpha-thalassemia.

**Family History**The patient's family history revealed no known autoimmune conditions, celiac disease, or inflammatory bowel disease among first-degree relatives. Genetic testing for her nonspecific genetic disorder provided inconclusive results.

**Initial Presentation of Illness**The patient reported chronic diarrhea for six months, characterized by loose, watery stools occurring six to seven times daily. This was accompanied by abdominal cramping, fatigue, and a marked reduction in body weight from 54 kg to 34 kg. She also reported intolerance to fatty foods and occasional bloating. No fever or vomiting was noted during this period.

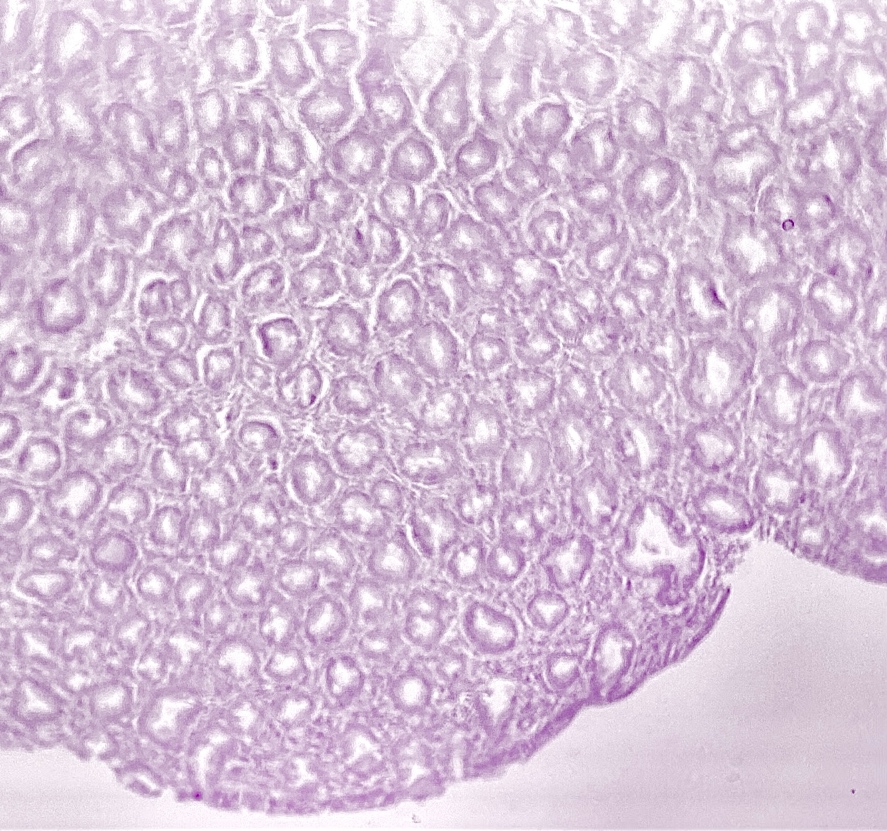
**Laboratory Investigations**  
Blood tests revealed microcytic hypochromic anemia (hemoglobin: 7.9 g/dL, normal: 12-15.5 g/dL; MCV: 76 fL, normal: 80-100 fL; MCHC: 29%, normal: 33-36%), hypoalbuminemia (28 g/L, normal: 35-50 g/L), hypocholesterolemia (0.8 g/L, normal: 1.6-2.5 g/L), and an elevated C-reactive protein (CRP) level of 66 mg/L (normal: <5 mg/L). Thrombocytosis (460,000/mm³, normal: 150,000-400,000/mm³) and ferritin elevation (250 ng/mL, normal: 30-200 ng/mL) were also observed. Liver function tests, renal function tests, and electrolyte levels were within normal ranges. Stool cultures and parasitology were negative for pathogenic organisms.

**Imaging and Endoscopy Findings**Colonoscopy revealed inflammatory changes in the right and left colon with ulceration in the cecum. The terminal ileum, transverse colon, and rectum appeared normal. Histopathological examination of biopsies showed non-specific colitis. Testing for Mycobacterium tuberculosis was negative.

Magnetic resonance imaging (MRI) showed terminal ileal thickening consistent with Crohn’s disease (Figure 1).

*Figure 1: Axial T1-weighted post-contrast MRI showing thickening of the terminal ileal loop (indicated by the white arrow).*

Upper gastrointestinal endoscopy demonstrated a mosaic pattern and total villous atrophy in the descending duodenum. Duodenal biopsies confirmed villous atrophy with intraepithelial lymphocytosis (40%), consistent with celiac disease. Immunology tests revealed elevated IgA anti-tissue transglutaminase antibodies (57 U/mL) and positive endomysial antibodies (1:302), confirming the CD diagnosis (figure 2).

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Figure 2: Image of microscopic examination (H&E stain, x25), revealing the presence of a total villous atrophy in the duodenal mucosa*

**Management and Follow-Up**The patient was initially treated with parenteral corticosteroids for five days, followed by oral corticosteroids and azathioprine. A strict gluten-free diet was initiated, resulting in symptomatic improvement and a 4 kg weight gain over three months.

**Outcome**   
Six months later, she presented with bilateral knee pain and morning stiffness lasting more than one hour. Rheumatological evaluation revealed peripheral arthritis and enthesitis. Imaging and clinical findings confirmed a diagnosis of ankylosing spondylitis (AS). Infliximab was added to her treatment regimen alongside azathioprine, leading to significant clinical remission. She regained an additional 6 kg over the following six months and reported no further gastrointestinal or joint symptoms.

3. discussion

Celiac disease (CD) and inflammatory bowel diseases (IBD) such as Crohn’s disease (CrD) are both chronic inflammatory conditions affecting the gastrointestinal tract [2]. Their co-occurrence was first described in 1967, but reports remain scarce in the literature [3]. Recent studies suggest an increasing incidence of IBD in patients with celiac disease. Bardella et al. demonstrated that patients with CD are eight times more likely to develop IBD compared to the general population [4,5].

Diagnosing both conditions in the same patient can be challenging, particularly when symptoms overlap. In Crohn’s disease, upper endoscopy may reveal duodenal scalloping, a feature typically associated with celiac disease due to villous atrophy [6].   
Moreover, anti-tissue transglutaminase antibodies (IgA-tTG) may occasionally be elevated in IBD patients, complicating the diagnosis.

Despite these diagnostic challenges, the presence of endomysial antibodies and significant intraepithelial lymphocytosis (≥30%) remains specific for celiac disease.

Pathophysiologically, both conditions share autoimmunological features, and both are associated with other autoimmune diseases, such as thyroiditis and vitiligo.   
Genetic predisposition has also been suggested, with familial forms observed in both conditions.   
A 2011 systematic review identified four susceptibility genes common to both CD and CrD: IL18RAP (interleukin-18 receptor accessory protein in 2q12), PUS10 (pseudouridylate synthase 10 in 2p16), TAGAP (T-cell activation Rho GTPase activating protein in 6q25), and PTPN2 (tyrosine-protein phosphatase non-receptor type 2 in 18p11) [7].

Latiano et al. also highlighted the role of the MYO9B gene, typically associated with CD, in IBD patients [8].

Finally, the association between IBD and ankylosing spondylitis (AS) is well documented, with AS affecting 2-3% of IBD patients, particularly those with CrD. The introduction of biologic therapies, specifically anti-TNF agents like infliximab [9] has revolutionized treatment for patients with AS and concomitant IBD, as seen in our case.

This highlights the importance of comprehensive treatment strategies in managing complex cases of overlapping autoimmune disorders.

4. Conclusion

The coexistence of Crohn’s disease and celiac disease is rare, with most cases of Crohn’s disease occurring after a pre-existing diagnosis of celiac disease. In this case, both conditions were diagnosed concurrently during the same hospitalization. While the association between ankylosing spondylitis and IBD is well-documented, to our knowledge, this is the first report describing the coexistence of Crohn’s disease, celiac disease, and ankylosing spondylitis in a single patient.

Consent

All authors declare that ‘written informed consent was obtained from the patient for publication of this case report and accompanying images.

ethical approval

Not applicable

Competing interests

Authors have declared that no competing interests exist

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